PET Radiopharmaceuticals for Imaging Chemotherapy-Induced Cardiotoxicity

Jothilingam Sivapackiam¹ · Monica Sharma¹ · Thomas H. Schindler¹,² · Vijay Sharma¹,³,⁴

Published online: 19 June 2020 © The Author(s) 2020

Abstract
Purpose of Review Currently, cardiotoxicity is monitored through echocardiography or multigated acquisition scanning and is defined as 10% or higher LVEF reduction. The latter stage may represent irreversible myocardium injury and limits modification of therapeutic paradigms at earliest stages. To stratify patients for anthracycline-related heart failure, highly sensitive and molecularly specific probes capable of interrogating cardiac damage at the subcellular levels have been sought.

Recent Findings PET tracers may provide noninvasive assessment of earliest changes within myocardium. These tracers are at nascent stages of development and belong primarily to (a) mitochondrial potential-targeted and (b) general ROS (reactive oxygen species)-targeted radiotracers. Given that electrochemical gradient changes at the mitochondrial membrane represent an upstream, and earliest event before triggering the production of the ROS and caspase activity in a biochemical cascade, the former category might offer interrogation of cardiotoxicity at earliest stages exemplified by PET imaging, using ¹⁸F-Mitophos and ⁶⁸Ga-Galmydar in rodent models.

Summary Both categories of radiotracers may provide tools for monitoring chemotherapy-induced cardiotoxicity and interrogating therapeutic efficacy of cardio-protectants.

Keywords Cardiotoxicity · PET radiopharmaceuticals · Anthracycline · ABCB1 · ABCG2 · Galmydar

Introduction

Over the last decade, successes in chemotherapy have contributed enormously to improvements in survival rates of cancer patients. During this intermediate period, the design and development of targeted therapeutics and characterization of novel biomarkers that mediate the pathophysiology of tumor biology in general, and their distant metastases in particular, have both demonstrated unparalleled growth, thereby advancing substantially the fields of medicinal chemistry, chemical, and tumor biology. Among these discoveries, while therapeutics directed at signaling pathways (angiogenesis) to impede proliferation of tumor cells via kinase inhibitors have shown promising outcomes, the discovery of immune check-point inhibitors (ICIs) have also offered remarkable tools in kits of molecular oncologists to treat both solid- and hematological tumors. Among these ICIs, agents targeting cytotoxic T lymphocyte-associated antigen 4 (CTLA-4), programmed cell death receptor-1 (PD-1), and programmed cell death ligand-1 (PD-L1) have found remarkable clinical success [1, 2]. The mechanism of action for these agents involve inhibiting the tumor cells from inactivating the immune system, thereby...
restoring the immune system role against the infiltrating tumor cells, thus impacting positively the healthcare landscape by enhancing survival rates in patient populations with typically poor outcomes using other treatment paradigms [3, 4]. However, each therapeutic advance has encountered its own challenges; therefore, ICIs also could not escape that fate. Importantly, the stimulation of the immune system is not without risk and is known to be associated with multi-organ adverse events [5]. These adverse events have been shown to occur with a variable frequency depending on the type of ICI, the type and location of the tumor, and host-traits [6]. Despite these advances, anthracyclines (exemplified by Doxorubicin or related analogues) either as primary intervention or as a combination therapy continue to be the main workhorse of patient management plans in molecular oncology clinics due to their utility in numerous malignancies (but not limited to), such as acute leukemia, non-Hodgkin lymphoma, breast cancer, and sarcomas. However, the benefits of these advances in cancer chemotherapy have been offset by concerns about cardiotoxicity-related adverse effects induced by administered anthracyclines or combination therapy. Most chemotherapeutic agents damaging myocardium often affect the circulatory vessels, impair coronary endothelial function, and can potentially induce left ventricular dysfunction or heart failure by generation of reactive oxygen species, and apoptosis [7]. Cardiotoxicity may be acute, which occurs either during or immediately following treatment and can be either transient or chronic, and has been typically categorized into type I (early onset) and type II (late onset) [8]. While the type I cardiotoxicity has been considered to be irreversible cardiac cell injury, and normally caused by anthracyclines, and analogues thereof including combination therapy, the type II cardiotoxicity is typically induced by molecularly targeted antibodies [9].

Cardio-oncology is a rapidly emerging field for managing cancer-therapy-related cardiotoxicity, the consensus around using left ventricle ejection fraction (LVEF) is one of the metrics, and is evolving criterion. For example, the Cardiac Review and Evaluation Committee of trastuzumab-associated cardiotoxicity defines it by symptoms of heart failure, decline of LVEF, either as symptomatic LVEF decrease from ≥5 to < 55% or an asymptomatic reduction of LVEF ≥ 10 to < 55% [10]. The American Society of Echocardiography and European Association of Cardiovascular Imaging define cardiotoxicity as global longitudinal strain (GLS) with a 10–15% early reduction. Finally, FDA defines LVEF drop < 40–45% or is 40–49% with a ≥10% absolute decrease below baseline with anti-HER2 targeted therapy as a benchmark for monitoring cardiotoxicity [11]. While the consensus among clinical community would continue to emerge on defining features of cardiotoxicity, molecular imaging agents capable of offering early detection of cardiotoxicity in vivo could offer powerful noninvasive tools for understanding role of different biomarkers regulating its pathophysiology, while simultaneously allowing opportunities for stratification of therapeutic choices for guiding patient management in the twenty-first century.

**Mechanisms and Biomarkers**

While mechanisms of action for anthracyclines are multifactorial, the consensus among scientific community is that chemotherapy can potentially trigger increased production of reactive oxygen and reactive nitrogen species, lipid peroxidation, inflammation, induced cardiomyocyte apoptosis, interstitial fibrosis, and abnormal signaling of epidermal growth factor as well as β-arrestin (the highly conserved family of cytosolic adaptor proteins that contribute to many immune functions by orchestrating the desensitization and internalization of cell-surface G protein-coupled receptors (GPCRs) via well-studied canonical interactions), inhibition of nuclear topoisomerase II β, induced DNA damage, inhibition of vascular endothelial growth factor signaling, defective mitochondrial biogenesis, and calcium overloading [12–14]. Among the intracellular organelles, while maintaining their role as energy powerhouse, mitochondria have an essential role in myocardial tissue homeostasis; thus, either deterioration or partial impairment in their normal function leads to cardiomyocyte and endothelial cell death, consequently thus inducing cardiovascular dysfunction [15]. For example, the antiretroviral nucleoside reverse transcriptase inhibitors such as zidovudine may cause cardiac mitochondrial dysfunction through inhibition of DNA polymerase-gamma and induction of mitochondrial DNA mutations, thus leading to cardiomyopathy [16]. Overall, cardiac adverse effects can primarily be classified into 2 categories: (a) functional and (b) structural effects. Of note, seriously altered function may be completely dissociated from the structural effect, especially at an early stage [17]. Other than the functional deterioration, anthracyclines have also been shown to damage several proteins regulating cardiac muscle contractility including titin, the myofilament forming protein that regulates cardiac function leading to systolic and diastolic dysfunction [18]. Finally, the inter-individual variability in the susceptibility to chronic anthracycline-induced cardiotoxicity has also been reported. Therefore, the genetic variants with occurrence of drug-induced cardiotoxicity have also been identified, thus suggesting a potential role for gene polymorphisms that may control the metabolism of anthracyclines [19], detoxification of free radicals, and alterations in physiological iron levels. Finally, SNPs in Her2/neu Pro 1170 Ala polymorphism have also been identified in a subset of patients with increased risk of cardiotoxicity from trastuzumab therapy and postulated to be deployed with other risk biomarkers for stratification of patients [20]. These factors suggest challenges in encountering cardiotoxicity, while
presenting opportunities to design new probes potentially capable of monitoring noninvasively cardiac dysfunction to manage cancer treatment paradigms in twenty-first century.

Currently, commonly used noninvasive diagnostic biomarkers to assess anthracycline-induced cardiotoxicity are cardiac troponins, brain natriuretic peptide, and N-terminal fragment of natriuretic peptide. Of note, persistent elevation of cardiac troponin I levels post 1-month treatment of anthracycline has also led to prevalence of more cardiac adverse events in 84% patients at 3 years compared with patients showing normal levels of cardiac troponin I levels. Additionally, circulating microRNAs also offer promising noninvasive tools and have been evaluated in children and young adults treated with anthracycline chemotherapy [21]. Finally, other less commonly used biomarkers for cardiotoxicity include cytostatin C, galectin-3, interleukin 6, tumor necrosis factor α (TNFα), myeloperoxidase, and C-reactive protein [22]. However, due to their potential for being influenced by numerous micro-environmental factors and non-cardiovascular diseases, these circulating biomarkers lack desired sensitivity and disease specificity.

To interrogate cardiotoxicity at a molecular level, molecular imaging enables visualization, characterization, and quantification of biomarkers or physiological processes at cellular and subcellular levels in vivo. Therefore, it is conceivable that cardiac nuclear imaging agents (PET and SPECT) may provide versatile diagnostic tools to detect cardiotoxicity at earlier stages, while enabling interrogation of therapeutic efficacy to afford stratification of chemotherapeutic choices in molecular oncology for better management of cardiotoxicity concerns.

**Nuclear Imaging Tracers**

Currently, cardiotoxicity is evaluated by echocardiography or multigated acquisition scanning and is typically defined as 10% or higher reduction of LVEF. However, a loss of such a magnitude in contraction function can be an indication of significant irreversible myocardium injury, thus substantially diminishing opportunities for interventions or modification of treatment plans. Therefore, interrogation of myocardium abnormalities at subcellular level may provide early and sensitive readout of drug-induced cardiotoxicity. Although pathophysiology of chemotherapy induced heart failure is a complex phenomenon involving multiple intersecting biochemical pathways, anthracycline-induced effects at a subcellular level have been known to be attributed to mitochondrial dysfunction or partial impairment in its normal function and elevated levels of oxidative stress. Arguably, nuclear imaging tracers have been under development based upon both themes. While the first category of tracers find their roots in application of fluorescent lipophilic cations and their utility in reporting about alterations in mitochondrial potential in cells [23], the second category of tracers has focused upon exploiting utility of dihydroethidium derivatives capable of detecting ROS in vivo. While the fluorescent lipophilic cations such as tetramethylrhodamine ethyl ester (TMRE) and Rhodamine 123 lack the desired depth penetration for whole body scanning, 99mTc-Sestamibi, a hydrophobic and monocationic technetium(I) octahedral complex, has been routinely used in clinic as a myocardial perfusion imaging agent. Importantly, 99mTc-Sestamibi also enables interrogation of mitochondrial potential similar to that of TMRE and Rhodamine 123 recently has been used to evaluate anthracycline-induced cardiotoxicity [24] in rodents. However, myocardium retention may need to be corrected using either pharmacokinetic modeling or normalized to another FDA-approved tracer to determine net signal in myocytes for diagnosis of cardiotoxicity. Compared with SPECT, PET tracers provide high sensitivity and enable quantification, thus allowing 3-dimensional pharmacokinetic analysis. Therefore, taking advantage of principles of lipophilic cations, 18F-Mitophos, a triphenylphosphonium analogue, has been recently evaluated for its potential to image DOX-induced cardiotoxicity in rat models [25]. The agent may be susceptible to concerns of perfusion effects similar to 99mTc-Sestamibi and also indicates significant metabolism, thus indicating reduction of parental tracer within minute post administration in vivo. Furthermore, 18F-DHMT, a dihydroethidium derivative [26, 27], has been also evaluated to monitor anthracycline-induced ROS cardiotoxicity in vivo [28]. Compared with significant effects observed in LVEF at 6-week post DOX-treatment, 18F-DHMT enables detection of superoxide production at 4-week post treatment [28]. Following further validations, the PET agent may allow opportunities for quantifying therapeutic efficacy of cardioprotectants, such as Dexrazoxane, an FDA-approved drug for late stage treatment of breast cancer patients [29].

While 99mTc-incorporated radiotracers have been the workhorse in nuclear medicine for decades due to commercial availability of 99Mo/99mTc generators in nuclear pharmacies, however, disruptions in the supply chain during the 99Mo/99mTc crisis of 2008–2010 demonstrated the vulnerability of the world supply of 99Mo. To address these shortcomings, over the last decade, germanium/gallium (Ge/Ga) generators have become available and are capable of producing high quality 68Ga (t1/2 = 68 min), an isotope with excellent emission properties for clinical PET imaging [30]. The parent isotope, 68Ge (t1/2 = 271 days), is produced in high-energy proton accelerators from a 68Ga(p,2n)68Ge reaction and is bonded to alumina for eventual elution on-site [31], thus providing a practical generator-based distribution model for on-site formulation of PET radiopharmaceuticals. With co-development of high-quality 68Ga-based tracers, PET imaging could be unlinked from proximity to cyclotrons, thereby expanding access to the technology. In next section, we
discuss development of another molecular imaging agent, described in literature as $^{68}$Ga-Galmydar, to track anthracycline induced effects in cellulo and in vivo.

Importantly, $^{68}$Ga-Galmydar is also recognized by ATP-binding-cassette (ABC) family of transporters, such as ABCB1 (also known as P-glycoprotein, 170 kD protein located on plasma membrane of tumor cells) and ABCG2 (also known as breast cancer resistance protein, BCRP, 72 kD protein) as their transport substrate [32]. Overall, the net retention of $^{68}$Ga-Galmydar in heart tissue is determined by the opposing action of two biochemical processes [33]. Deploying delocalization of charge on its molecular surface, this tracer permeates passively into living cells and concentrating within the mitochondrial inner matrix in response to the driving forces of electronegative plasma membrane and mitochondrial transmembrane potentials [34]. However, this uptake is opposed by action of ABC membrane transporters, such as ABCB1, MRP1 (ABC11), and ABCG2, which excrete the radiotracer into extracellular space of tumor cells [35–39]. Importantly, cardiomyocytes, although rich in mitochondria, they lack expression of efflux transporter proteins, thus sequestering this radiotracer for prolonged periods to enable imaging, while hepatocytes, which express ABCB1 and ABCG2 along their cannalicular surface, rapidly excrete it into the bile and intestines. In principle, these biochemical traits would be expected to facilitate hepatocellular clearance of the tracer, thereby minimizing the impact of $\gamma$-emissions arising from the liver that could potentially co-register into the inferior wall of the myocardium during imaging.

Galmydar is also a mildly fluorescent molecular imaging probe and localizes within the mitochondria of rat cardiomyoblasts (H9c2) similar to mitotracker Red [40–42] (Fig. 1). While tracer shows stable accumulation in rat cardiomyoblasts, its uptake profiles in MCF-7 cells are inversely proportional to expression of ABCB1 on the plasma membrane (Fig. 2) [39]. Similar uptake profiles are observed...
in HEK 293 ABCG2 transfected cells [32, 43, 44]. The agent shows dose and time-dependent pharmacological response to anthracycline in rat cardiomyoblasts [45••], using live cell imaging, thereby consistent with postulated mechanism of anthracycline-induced depolarization of mitochondrial redox potentials, and ROS production [46]. To further assess potential of 

$^{68}$Ga-Galmydar to serve as a molecular imaging probe for assessment of DOX-induced cardiotoxicity in vivo, $^{68}$Ga-Galmydar has been injected via tail-vein into rats pretreated either with intravenous administration of DOX (15 mg/kg) or vehicle (5% ethanol in saline). Following treatments, micro-PET static scans (10-min acquisition; 60-min post tail-vein administration of $^{68}$Ga-Galmydar; Fig. 3a) demonstrated a 1.91-fold lower retention in hearts of DOX-treated (Standard Uptake Value; SUV: 0.92, $n=3$) rats compared with their vehicle-treated counterparts (SUV: 1.76, $n=3$; Fig. 3b) [45••]. For correlation of PET data, post-imaging quantitative biodistribution studies demonstrate heart retention values of 2.02-folds lower for DOX treated (%ID/g; DOX: 0.44 ± 0.1, $n=3$) rats compared to their vehicle-treated counterparts (vehicle control: 0.89 ± 0.03, $n=3$, $p=0.04$; Fig. 3c), thus supporting micro-PET imaging data in vivo [45••]. Of note, using live-cell imaging, Galmydar also indicate a gradual depression in cellular uptake and retention of Galmydar (up to 8.2-fold difference compared to their untreated cells after 5 h), thus indicating the sensitivity of the probe to map changes at the level of the mitochondria resulting from DOX treatment, which in turn likely result from depolarization of the mitochondrial potential [45••]. These findings are consistent with literature precedents, wherein DOX treatment has been shown to alter mitochondrial redox potentials, thus depolarizing mitochondria, and elevating matrix Ca2+ and ROS production in 30 min [46]. Finally, high-resolution single-cell imaging also shows localization of Galmydar in mitochondria of DOX-treated cells (Fig. 1, lower panel) similar to their untreated controls (top panel), however, substantial decreased retention thus correlating with the lower PET signal in heart of DOX-treated rats (Fig. 3). Following further validations in higher vertebrates, $^{68}$Ga-Galmydar imaging could enable monitoring of impaired mitochondrial function in myocytes following anthracycline treatment in vivo.

### Conclusions

While a significant loss in contractile function of the myocardium may serve as a warning for irreversible tissue damage, current imaging techniques may not have the desired sensitivity and molecular specificity to guide interventions at early stages of cardiotoxicity. Among various imaging modalities, nuclear imaging-based strategies can potentially be translated faster into clinic due to the need for administration of doses at very low concentrations. Both mitochondrial potential- and ROS-targeted tracers may allow noninvasive imaging of anthracycline-induced cardiotoxicity in vivo. Because frontiers of molecular imaging in twenty-first century are pushing the edge of the envelop to detection at earliest stages, it may be argued biochemically that changes in the mitochondrial potentials represent an upstream event, before triggering the production of the ROS and caspase activity; thus, it is conceivable that tracers capable of reporting changes in the mitochondrial potential in vivo might offer interrogation of cardiotoxicity at earliest stages as evident from imaging of $^{18}$F-Mitophos and $^{68}$Ga-Galmydar in rodent models. It remains to be determined, whether these initial observations would replicate in higher vertebrates and translate into humans. We envision that both categories of radiotracers could be beneficial for monitoring cardiotoxicity in the field of cardio-oncology and may provide opportunities for interrogating therapeutic efficacy of cardio-

![Fig. 3](image-url)
protectants, while offering opportunities for stratification of cancer patients for modification of therapeutic protocols.

Acknowledgments Authors are grateful to colleagues and coworkers (both current and past) within molecular imaging center and Sharma laboratory for their valuable suggestions and thoughtful contributions.

Funding Information This work was supported in part by grants RO1 HL111163 (VS) and RO1 HL142297 (VS) from the National Institutes of Health and departmental funds.

Compliance with Ethical Standards

Conflict of Interest Jothilingam Sivapackiam and Vijay Sharma are inventors of Galmydar (US 9,579,408; Washington University has IP rights; there is no licensee or royalties) and declare no competing financial interests. Monica Sharma and Thomas H. Schindler declare that they have no conflict of interest.

Human and Animal Rights and Informed Consent This article does not contain any new study and is a simple illustration of scientific information available in the literature.

Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit http://creativecommons.org/licenses/by/4.0/.

References

Papers of particular interest, published recently, have been highlighted as:

• Of importance
•• Of major importance

1. Sharma P, Allison JP. The future of immune checkpoint therapy. Science. 2015;348(6230):56–61.
2. Goodman A, Patel SP, Kurzrock R. PD-1-PD-L1 immune-checkpoint blockade in B-cell lymphomas. Nat Rev Clin Oncol. 2017;14(4):203–20.
3. Pitt JM, Vetizou M, Daillere R, Roberti MP, Yamazaki T, Routi B, et al. Resistance mechanisms to immune-checkpoint blockade in cancer: tumor-intrinsic and -extrinsic factors. Immunity. 2016;44(6):1255–69.
4. Hurst JH. Cancer immunotherapy innovator James Allison receives the 2015 Lasker–DeBakey Clinical Medical Research Award. J Clin Invest. 2015;125(10):3732–6.
5. Hofmann L, Forschner A, Loquai C, Goldinger SM, Zimmer L, Ugurel S, et al. Cutaneous, gastrointestinal, hepatic, endocrine, and renal side-effects of anti-PD-1 therapy. Eur J Cancer. 2016;60:190–209.
6. Hananishi J, Mandai M, Ikeda T, Minami M, Kawaguchi A, Murayama T, et al. Safety and antitumor activity of anti-PD-1 antibody, Nivolumab, in patients with platinum-resistant ovarian cancer. J Clin Oncol. 2015;33(34):4015–22.
7. Aggarwal S, Kamboj J, Arora R. Chemotherapy-related cardiotoxicity. Ther Adv Cardiovasc Dis. 2013;7(2):87–98.
8. Jain D, Russell RR, Schwartz RG, Panjehpour GS, Aronow W. Cardiac complications of cancer therapy: pathophysiology, identification, prevention, treatment, and future directions. Curr Cardiol Rep. 2017;19(5):36.
9. Tocchetti CG, Cadeddu C, Di Lisi D, Femminino S, Madonna R, Mele D, et al. From molecular mechanisms to clinical management of antineoplastic drug-induced cardiovascular toxicity: a translational overview. Antioxid Redox Signal. 2019;30(18):2110–53 This article outlines mitochondrion as a critical target for chemotherapy-induced cardiovascular toxicity, while authors also envision genetic profiling with molecular imaging could be potentially deployed for stratification of a risk factor for cardiotoxicity in cancer patients.
10. Seidman A, Hudis C, Pierri MK, Shak S, Paton V, Ashby M, et al. Cardiac dysfunction in the trastuzumab clinical trials experience. J Clin Oncol. 2002;20(5):1215–21.
11. Ponde NF, Lambertini M, de Azambuja E. Twenty years of anti-HER2 therapy-associated cardiotoxicity. ESMO Open. 2016;1(4): e000073 This is an important review article on anti-HER2 therapy, and other anti-HER2 specific drugs.
12. Gu J, Hu W, Zhang DD. Resveratrol, a polyphenol phytoalexin, protects against doxorubicin-induced cardiotoxicity. J Cell Mol Med. 2015;19(10):2324–8.
13. Novo G, Cadeddu C, Sucato V, Pagliaro P, Romano S, Tocchetti CG, et al. Role of biomarkers in monitoring antiinflammatory cardiotoxicity. J Cardiovasc Med (Hagerstown). 2016;17(Suppl 1):S27–34.
14. Varricchi G, Ameri P, Cadeddu C, Ghigo A, Madonna R, Marone G, et al. Antineoplastic drug-induced cardiotoxicity: a redox perspective. Front Physiol. 2018;9:167.
15. Varga ZV, Ferdinandy P, Liao L, Pacher P. Drug-induced mitochondrial dysfunction and cardiotoxicity. Am J Physiol Heart Circ Physiol. 2015;309(9):H1453–67.
16. Lewis W. Cardiomyopathy, nucleoside reverse transcriptase inhibitors and mitochondria are linked through AIDS and its therapy. Mitochondrion. 2004;4(2–3):141–52.
17. O’Brien PJ. Cardiac troponin is the most effective translational safety biomarker for myocardial injury in cardiotoxicity. Toxicology. 2008;245(3):206–18.
18. Sadurska E. Current views on anthracycline cardiotoxicity in childhood Cancer survivors. Pediatr Cardiol. 2015;36(6):1112–9.
19. Blanco JG, Sun CL, Landier W, Chen L, Esparza-Duran D, Leisenring W, et al. Anthracycline-related cardiotoxicity in children: role of polymorphisms in carbonyl reductase genes—a report from the Children’s Oncology Group. J Clin Oncol. 2012;30(13):1415–21.
20. Stanton SE, Ward MM, Christos P, Sanford R, Lam C, Cobham MV, et al. Pro1170 Ala polymorphism in HER2-neu is associated with risk of trastuzumab cardiotoxicity. BMC Cancer. 2015;15:267.
21. Riddell E, Lenihan D. The role of cardiac biomarkers in cardiology. Curr Probl Cancer. 2018;42(4):375–85 This is an interesting summary of vital biomarkers as risk factors for cardiotoxicity including microRNAs and myeloperoxidase as exploratory tools for identifying cardiac dysfunction.
22. Tian S, Hirshfield KM, Jabbour SK, Toppmeyer D, Haffty BG, Khan AJ, et al. Serum biomarkers for the detection of cardiac toxicity after chemotherapy and radiation therapy in breast cancer patients. Front Oncol. 2014;4:277.
23. Cottet-Rousselle C, Ronot X, Leverve X, Mayol JF. Cytometric assessment of mitochondria using fluorescent probes. Cytometry A. 2011;79(6):405–25.

24. Safaez ZM, Baark F, Waters ECT, Veronese M, Pell VR, Clark JE, et al. Detection of anthracycline-induced cardiotoxicity using perfusion-corrected (99mTc) sestamibi SPECT. Sci Rep. 2019;9(1):216.

25. McCluskey SP, Haslop A, Coelho C, Gunn RN, Tate EW, Southworth R, et al. Imaging of chemotherapy-induced acute cardiotoxicity with (18)F-labeled lipophilic cations. J Nucl Med. 2019;60(12):1750–6 This article describes ability of (18)F-Mitosphos to detect anthracycline-induced cardiotoxicity in rat models. Importantly, this agent is sensitive to detection of prevailing electrochemical gradients at the mitochondrial membrane. If indeed these data can be replicated in higher vertebrates and translated into humans, it may be able to detect earlier stages of anthracycline-induced cardiotoxicity to afford opportunities for modification of therapeutic protocols in molecular oncology clinic.

26. Chu W, Chepetan A, Zhou D, Shoghi KI, Xu J, Dugan LL, et al. Development of a PET radiotracer for non-invasive imaging of the reactive oxygen species, superoxide, in vivo. Org Biomol Chem. 2014;12(25):4421–31 This article is an original report of an (18)F-DHMT tracer for detection of ROS following anthracycline treatment.

27. Zhang W, Cai Z, Li L, Ropchan J, Lim K, Boutagy NE, et al. Optimized and automated radiosynthesis of [(18)F]DHMT for translational imaging of reactive oxygen species with positron emission tomography. Molecules. 2016;21(12):1696.

28. Boutagy NE, Wu J, Cai Z, Zhang W, Booth CJ, Kyriakides TC, et al. In vivo reactive oxygen species detection with a novel positron emission tomography tracer, (18)F-DHMT, allows for early detection of anthracycline-induced cardiotoxicity in rodents. JACC Basic Transl Sci. 2018;3(3):378–90. This is an important and robust PET imaging study showing ability of the radiotracer to detect ROS activity in rats 5–6 week post anthracycline treatments. The authors demonstrate that drop in LVEF is associated with significant increase in MMP activation, cellular apoptosis, and myocardial toxicity.

29. Kim IH, Lee JE, Youn HJ, Song BJ, Chae BJ. Cardioprotective effect of dexrazoxane in patients with HER2-positive breast cancer who receive anthracycline-based adjuvant chemotherapy followed by trastuzumab. J Breast Cancer. 2017;20(1):82–90.

30. Pagou M, Zervier I, Al-Nahas A. Can gallium-68 compounds partly replace (18)F-FDG in PET molecular imaging? Hell J Nucl Med. 2009;12(2):102–5.

31. Fani M, Andre JP, Maecke HR. 68Ga-PET: a powerful generator-based alternative to cyclotron-based PET radiopharmaceuticals. Contrast Media Mol Imaging. 2008;3(2):67–77.

32. Sivapackiam J, Harpstrite SE, Prior JL, Mattingly S, Sharma V. (67/68)Galmymdar: a metalloprotease for monitoring breast cancer resistance protein (BCRP)-mediated functional transport activity. Nucl Med Biol. 2016;43(3):191–7.

33. Sharma V, Beatty A, Wey S-P, Dahlheimer J, Pica C, Crankshaw C, et al. Novel gallium(III) complexes transported by MDR1 P-glycoprotein: potential PET imaging agents for probing P-glycoprotein-mediated transport activity in vivo. Chem Biol. 2000;7:335–43.

34. Davis S, Weiss M, Wong J, Lampidis T, Chen L. Mitochondrial and plasma-membrane potentials cause unusual accumulation and retention of rhodamine-123 by human breast adenocarcinoma-derived MCF-7 cells. J Biol Chem. 1985;260:13844–50.

35. Hendriks N, de Vries EGE, Eriks-Fluks L, van der Graaf W, Hospers G, Willemsen A, et al. A new in vivo method to study P-glycoprotein transport in tumors and the blood-brain barrier. Cancer Res. 1999;59:2411–6.

36. Hendriks N, Franssen E, van der Graaf W, Meijer C, Piers D, Vaalburg W, et al. 99mTc- Sestamibi is a substrate for P-glycoprotein and the multidrug resistance-associated protein. Br J Cancer. 1998;77:353–8.

37. Hendriks N, Franssen E, van der Graaf W, de Vries E. Visualization of multidrug resistance in vivo. Eur J Nucl Med. 1999;26:283–93.

38. Sharma V. Radiopharmaceuticals for assessment of multidrug resistance P-glycoprotein-mediated transport activity. Bioconjug Chem. 2004;15:1464–74.

39. Sharma V, Sivapackiam J, Harpstrite SE, Prior JL, Gu R, Rath NP, et al. A generator-produced gallium-68 radiopharmaceutical for PET imaging of myocardial perfusion. PLoS One. 2014;9(10):e109361.

40. Takayama S, Ostuni E, LeDuc P, Naruse K, Ingber DE, Whitesides GM. Subcellular positioning of small molecules. Nature. 2001;411(6841):1016.

41. Vilas GL, Corvi MM, Plummer GJ, Seime AM, Lambkin GR, Berthiaume LG. Posttranslational myristoylation of caspase-activated p21-activated protein kinase 2 (PAK2) potentiates late apoptotic events. Proc Natl Acad Sci U S A. 2006;103(17):6542–7.

42. Shrivastava A, Tiwari M, Sinha RA, Kumar A, Balapure AK, Bajpai VK, et al. Molecular iodine induces caspase-independent apoptosis in human breast carcinoma cells involving the mitochondria-mediated pathway. J Biol Chem. 2006;281(28):19762–71.

43. Doyle L, Ross DD. Multidrug resistance mediated by the breast cancer resistance protein BCRP (ABCG2). Oncogene. 2003;22(47):7340–8.

44. Zhou S, Schuetz JD, Bunting KD, Colapietro AM, Sampath J, Morris JJ, et al. The ABC transporter Bcrp1/ABCG2 is expressed in a wide variety of stem cells and is a molecular determinant of the side-population phenotype. Nat Med. 2001;7(9):1028–34.

45. Sivapackiam J, Kabra S, Speidel S, Sharma M, Lafrost R, Salter A, et al. 68Ga-Galmymdar: a PET imaging tracer for noninvasive detection of Doxorubicin-induced cardiotoxicity. PLoS One 2019;14(5):e0215579. https://doi.org/10.1371/journal.pone.0215579. eCollection 2019. This is an important contribution to the field, wherein cardiotoxicity is evaluated using both live-cell fluorescence imaging and PET/CT imaging deploying 68Ga-Galmymdar thus correlating information from PET imaging to that of optical imaging at a single cell level.

46. Lu P. Monitoring cardiac function in patients receiving doxorubicin. Semin Nucl Med. 2005;35(3):197–201.

Publisher’s Note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.