

Ferumoxytol

Since the Food and Drug Administration ([FDA](#)) approved ferumoxytol (Feraheme, AMAG Pharmaceuticals, Waltham, MA) to treat iron deficiency anemia in adults with chronic kidney disease (CKD) in 2009, the off label use of this iron oxide nanoparticle compound by clinicians and researchers as a magnetic resonance imaging (MRI) contrast agent has grown rapidly. Ferumoxytol-enhanced imaging is feasible in patients with impaired renal function, a patient population in whom both gadolinium and iodinated contrast agents are contraindicated. Other attractive imaging features of intravenous (IV) ferumoxytol include a prolonged blood pool phase and delayed intracellular uptake. Furthermore, since iron is a naturally occurring element in the body, the administered iron enters the body's natural iron metabolic pathways. Thus, the use of ferumoxytol is not currently associated with concerns regarding long-term deposition, as is the case with brain deposition of gadolinium-containing agents ^{1) 2) 3) 4)}

see [Ferumoxytol magnetic resonance imaging](#).

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