Precision in Musculoskeletal imaging: Utilizing Dixon method to Distinguish Non-Neoplastic from Neoplastic lesions

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Learning outcomes

- ✤ The composition and roles of red and yellow marrow
- ✤ Physiological differences between non-neoplastic and neoplastic lesions
- ✤ Signal Suppression techniques in MRI
- ✤ Limitations of routine MRI sequences in MSK imaging
- ✤ Physical principles of Dixon and its diagnostic accuracy in MSK imaging
- ✤ A case study



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Bone marrow development and contents

In the last 3-month before the birth and after birth, **Red marrow** plays a vital role in Haemopoiesis.

INTRAMEDULLARY COMPOSITION OF RED MARROW VS. YELLOW MARROW

Red marrow		Yellow marrow
Triglycerides (fat), 40%	intramedullary fluid, 40%	Triglycerides (fat), 80%
Haempoietic cell, 20%		intermedullary fluid, 20%



(Alexander, Laor and Bedoya, 2023)

(Jahanvi and Kelkar, 2021) (Omoumi, 2022)

Marrow Disorders



Benign neoplastic Aneurysmal bone cyst Atypical haemangioma Benign notochordal tumour Chondroblastoma Fibrous cortical defect Giant cell carcinoma (benign) Non-occyfying fibroma Osteoma Simple bone cyst Schwannoma Enchondroma

Malignant neoplastic Chondrosarcoma Chordoma Ewing sarcoma Giant cell carcinoma (malignant) Leukaemia **Metastasis** Myeloma Osteosarcoma Plasmacytoma Pleomorphic sarcoma Round cell carcinoma Spindle cell sarcoma

Malignant Neoplastic lesions

(Kohl et al., 2014) (Saifuddin et al., 2021) (Douis et al., 2016) (Van Vucht et al., 2021)

Routine musculoskeletal MRI sequences

In T1-W imaging, it is hard to precisely delineate **intramedullary borders of bone tumours** from extensive peritumoral inflammation or abundant red marrow. (Shiga et al. 2013)

In T1-W+Contrast imaging, the enhancement can be **exaggerated** by the inflammatory responses around the detected lesion.

(Verstraete and Lang, 2000)

Other fat-suppression techniques such as **STIR** and **SPAIR** cannot be used for contrast-enhancement imaging.

(Kenneally et al., 2015)



Coronal T1-Weighted of R.t femur



Axial T1-Weighted of R.t femur



Axial STIR of R.t femur

Physical Principles in Dixon Method



 $S_{OP} = Water_{signal} - Fat_{signal}$



 $S_{WO} = \frac{S_{IP} + S_{OP}}{2}$



 $S_{IP} = Water_{signal} + Fat_{signal}$







Percentage of Signal Intensity Drop (%SI Drop)

We searched seven academic databases:

- 1. Pub med
- 2. EBSChost
 - 1. CINHAL
 - 2. Academic Search Ultimate
 - 3. MEDLINE Complete
- 3. Ovid Library
 - 1. Ovid Emcare
 - 2. Embase
 - 3. Mediline

 $\% SI \ drop = \left[\frac{SI_{IP} - SI_{OP}}{SI^{IP}}\right] \times 100$

Magnetic field strength:	%SI Drop in Non- neoplastic lesions	%SI Drop in Neoplastic lesion			
1.5T	>20%	<20%			
3.0T	>25%	<25%			

We selected 11 articles (;4 primary+7 secondary research papers) based on inclusion & exclusion criteria.

Diagnostic value of %SI Drop in Dixon method

Study name	Object of study %	Sensitivity %	Specificity %	PPV %	NPV %	Accuracy %
	Using T1W GRE Dixon to distinguish non-neoplastic from neoplastic (benign + malignant) lesions	91.70	72.70	47.80	97.10	82.50
Douis et al., 2016 (n=57)	% SI drop (non-neoplastic)= 36.3%					
	% SI drop (malignant lesions) = 3.3-4.0%					
Kohl et al., 2014	To assess %SI drop between IP and OP images of T1W FSPGR to classify marrow lesions.	100	61	75	100	82
Saifuddin et al., 2021 (n=85)	Detecting non-neoplastic lesions with T1W GRE Dixon	66.70	88.10	61.9	90.9	84.70
	% SI drop (non-neoplastic)= 33.9-34.1%	72.20	05.10	50.2	02.1	02.50
	Detecting non-neoplastic lesions with T2W FSE Dixon	72.20	85.10	58.3	92.1	83.50
	% SI drop = 40.7-42.1%	72.20	00.00	CD D	02.2	04.70
	Detecting non-neoplastic lesions with T1W GRE Dixon	72.20	89.60	63.2	92.2	84.70
	% SI drop (benign lesions) = 7.8-12.7%					
	% SI drop (malignant lesions) = -9.0-4.0%					
	Detecting neoplastic (benign + malignant) lesions with T2W GRE Dixon	77.80	86.60	59.1	93.4	83.50
	% SI drop (benign lesions) = 14.4-15.4%					
	% SI drop (malignant lesions) = 3.3-4.0%					
Van Vucht et al., 2021 (n=174)	Using T1W GRE Dixon to distinguish non-neoplastic from neoplastic (benign + malignant) lesions	65.90	94.60	80.6	89.1	87.40
	% SI drop (non-neoplastic) = 36.6%					
	% SI drop (benign lesions) = 3.19%					
	% SI drop (malignant lesions) = 3.24%					
	Using T1W GRE Dixon to distinguish benign neoplastic from malignant neoplastic lesions	9.10	98.40	85.7	51.2	53.10

The pooled sensitivity, specificity, and accuracy of T1-W Dixon imaging to distinguish non-neoplastic from neoplastic (benign & malignant) lesions were 85.86%, 76.1%, and 83.96% respectively

Case Study

 $\%SI \, drop_{min} = \left[\frac{90.5 - 2.82}{90.5}\right] \times 100$ %*SI drop* = **96**.**88**% $\%SI \ drop_{mean} = \left[\frac{336.9 - 51.11}{336.9}\right] \times 100$ %*SI drop* = **84**. **83**% $\% SI \, drop_{max} = \left[\frac{550.36 - 298}{550.36}\right] \times 100$

%*SI drop* = **45**.**85**%

$\%SI \, drop = \left[\frac{SI_{IP} - SI_{OP}}{SI^{IP}}\right] \times 100$

%Signal Drop



Fibrous dysplasia; Non-neoplastic lesion

Conclusion

- When routine MRI sequences are used in conjunction with the Dixon method, the diagnostic capability of MRI to differentiate between non-neoplastic and neoplastic lesions is enhanced.
- The Dixon method provides multiple image sets (OP, IP, WO, and FO) within a single acquisition, more homogenous fat suppression, and allowing for both pre- and post-contrast imaging (; unlike other CSI techniques, such as STIR and SPAIR).
- Quantitative data derived from the Dixon method can facilitate more definitive interpretation of bone marrow lesions, thus improving patient management.
- The use of the Dixon method can reduce the need for additional invasive procedures, such as bone biopsies and contrast-enhanced imaging, particularly in cases of non-neoplastic lesions.
- However, bone biopsy may still be necessary to differentiate intermediate lesions (;malignant vs. benign neoplastic).

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THANKS!

Any questions?

The End



William Thomas Dixon (1945-2022)

Image caption: Tom Dixon receives the Gold Medal Award at the 2013 ISMRM Annual Meeting, Salt Lake City, Utah, USA.

Tom died on August 1, 2022, due to complications from Parkinson's disease.