

# **Original Research Article**

# DIAGNOSTIC EFFICACY OF O-RADS MRI IN DIFFERENTIATING BENIGN AND MALIGNANT ADNEXAL MASSES: A PROSPECTIVE STUDY WITH HISTOPATHOLOGICAL VALIDATION IN SOUTH INDIA

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## **ABSTRACT**

**Background:** Adnexal lesions are common in clinical practice, with a small but significant proportion being malignant. Ultrasound is the first-line tool; however, indeterminate lesions are not uncommon. The Ovarian-Adnexal Reporting and Data System for MRI (O-RADS MRI), introduced by the American College of Radiology (ACR) and ESUR, provides a standardized risk stratification system incorporating morphological features, diffusion-weighted imaging (DWI), and dynamic contrast enhancement (DCE). Several multicenter studies have validated its high diagnostic accuracy, but limited data are available from South Asia. The aim is to evaluate the diagnostic performance of O-RADS MRI in differentiating benign from malignant adnexal lesions, using histopathology as the reference standard. Materials and Methods: This prospective single-centre study was conducted between June 2024 and May 2025. A total of 73 women with adnexal masses underwent pelvic MRI including DWI and DCE sequences. Lesions were classified according to O-RADS MRI (categories 1-5). MRI-based categories were compared with histopathology, with premalignant and malignant lesions grouped together. Sensitivity, specificity, PPV, NPV, and diagnostic accuracy were calculated. Result: Of 73 adnexal lesions, 22 (30.1%) were malignant. O-RADS MRI showed a sensitivity of 93.1%, specificity of 87.5%, PPV 77.4%, NPV 96.4%, and overall accuracy of 89.9%. Conclusion: O-RADS MRI is a highly effective, standardized, and reproducible system for stratifying adnexal lesions. It demonstrates excellent diagnostic performance in distinguishing benign from malignant lesions and can play a critical role in guiding management, particularly in indeterminate cases after ultrasound.

### INTRODUCTION

Ovarian cancer is the fifth leading cause of cancer-related mortality among women worldwide, with late diagnosis contributing significantly to poor outcomes. [1,2] Accurate preoperative characterization of adnexal lesions is critical to avoid unnecessary surgeries for benign lesions and to ensure timely referral of malignant cases. [3,4] While ultrasound remains the first-line modality, indeterminate findings are not uncommon. [5] MRI, with its superior soft tissue contrast and functional sequences, plays an important role in further characterizing these lesions. [6] The O-RADS MRI scoring system was

developed to standardize reporting, provide objective criteria, and improve risk stratification.<sup>[7]</sup> This study aims to evaluate its diagnostic performance in an tertiary care setting in South India.

### **Review of Literature**

Several studies have validated O-RADS MRI's diagnostic performance. Thomassin-Naggara et al. demonstrated high sensitivity and specificity in differentiating benign from malignant adnexal masses. [8,9] Sadowski et al., in a multicenter study, reported sensitivity of 91.5% and specificity of 85%. [10] Forstner et al. confirmed reproducibility and clinical utility across European centers. [11] Vargas et al. emphasized the value of multiparametric MRI incorporating DWI and DCE. [12] Andreotti et al.

highlighted the role of O-RADS MRI in standardized risk stratification.<sup>[13]</sup> Despite robust international data, regional validation in South Asia remains limited.<sup>[14,15]</sup>

## MATERIALS AND METHODS

**Design:** Prospective observational study

**Period:** June 2024 – May 2025

Subjects: 73 women with adnexal lesions detected

on ultrasound and referred for MRI

# **Inclusion Criteria**

Age >18 years, adnexal lesion confirmed on MRI, histopathology available.

#### **Exclusion Criteria**

Known advanced gynecological malignancy, no histological follow-up.

MRI Protocol: 1.5T scanner; sequences included T1, T2, fat-suppressed T1, DWI (b=0, 800), ADC maps, and DCE perfusion.

**Interpretation:** Two radiologists independently applied O-RADS MRI scoring (v1.0, 2018). Discordances resolved by consensus.

**Histopathology:** Gold standard. Premalignant and malignant grouped together.

#### RESULTS

The commonest age group was 40–50 years (34.2%), followed by 50–60 years.

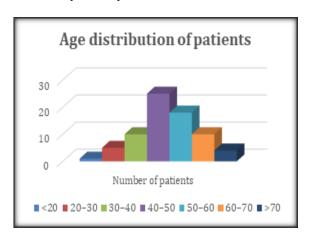
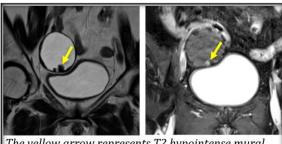


Table 1: O-RADS MRI category vs Histopathology outcome

O-RADS MRI Category	Benign (n)	Malignant (n)
2	37	1
3	11	2
4	4	7
5	1	12

Table 2: Diagnostic performance of O-RADS MRI

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Metric	Value	
Sensitivity	93.1%	
Specificity	87.5%	
Positive Predictive Value	77.4%	
Negative Predictive Value	96.4%	
Accuracy	89.9%	



The yellow arrow represents T2 hypointense mural nodule within the cyst on the left T2WI which shows post contrast enhancement on the right DCE image.

# **DISCUSSION**

Our study demonstrates that O-RADS MRI has excellent diagnostic performance in differentiating benign from malignant adnexal lesions, with sensitivity (93.1%), specificity (87.5%), and accuracy (89.9%) closely matching international data. Thomassin-Naggara et al. first validated the O-RADS MRI scoring system with sensitivity of 93% and specificity of 91%, [8] while Sadowski et al. reported similar values in a large multicenter trial. [10]

These results support the robustness and reproducibility of the system across different clinical settings.

Compared with ultrasound-based O-RADS, MRI offers superior tissue characterization, especially with diffusion-weighted imaging and dynamic contrast enhancement. [9,12] In our series, only one O-RADS 2 lesion was malignant, highlighting the system's high negative predictive value (96.4%), a key factor in ruling out malignancy and avoiding unnecessary surgery.

Our findings also align with Forstner et al,<sup>[11]</sup> who emphasized the utility of MRI in sonographically indeterminate adnexal masses, and Vargas et al,<sup>[12]</sup> who showed the added value of multiparametric imaging in refining risk stratification. Importantly, the standardized lexicon of O-RADS MRI enhances interobserver agreement and provides a reproducible framework for clinical decision-making.<sup>[7,13]</sup>

The main limitations of our study include its singlecenter design and modest sample size, similar to other regional studies.<sup>[14,15]</sup> Larger multicenter Indian data are needed to confirm external validity. Nevertheless, this prospective study provides important evidence supporting O-RADS MRI in South Asian populations, where data remain scarce.

#### CONCLUSION

O-RADS MRI is a robust and standardized system for adnexal lesion risk stratification, offering high diagnostic accuracy and reproducibility. By minimizing unnecessary surgeries for benign lesions and enabling early identification of malignant cases, it ensures timely and effective patient care. Wider implementation in India through clinician training, multidisciplinary awareness, and integration into national guidelines has the potential to significantly enhance early detection, streamline management pathways, and ultimately better survival outcomes.

### REFERENCES

- Karanth H, Murali S, Koteshwar R, Shetty V, Adappa K. Comparative study between propofol and dexmedetomidine for conscious sedation in patients undergoing outpatient colonoscopy. Anesthesia Essays and Researches. 2018 Jan 1;12(1):98 102.
- Mahadevan V. Anatomy of the caecum, appendix and colon. Surgery (Oxford). 2020 Jan 1;38(1):1 6.
- Boutros M, Gordon PH. Anatomy and physiology of the colon, rectum, and anal canal. Current therapy in colon and rectal surgery, 3rd edn. Elsevier, Philadelphia. 2017:3 11.
- Carrington EV, Scott SM. Physiology and function of the colon. Advanced Nutrition and Dietetics in Gastroenterology. 2014 Aug 25:28 32.
- Bhagatwala J, Singhal A, Aldrugh S, Sherid M, Sifuentes H, Sridhar S. Colonoscopy indications and contraindi cations. Screening for Colorectal Cancer with Colonoscopy. 2015 Dec 2:35 47.
- Rex DK, Schoenfeld PS, Cohen J, Pike IM, Adler DG, Fennerty BM, Lieb JG, Park WG, Rizk MK, Sawhney MS, Shaheen NJ. Quality indicators for colonoscopy. Official journal of the Amer ican College of Gastroenterology ACG. 2015 Jan 1;110(1):72 90.
- Baker FA, Mari A, Hosadurg D, Suki M, Ovadia B, Gal O, Kopelamn Y. The impact of colonoscopy indication on polyp detection rate. Annals of Gastroenterology. 2019 May;32(3):278.
- Dossa F, Mede iros B, Keng C, Acuna SA, Baxter NN. Propofol versus midazolam with or without short acting opioids for sedation in colonoscopy: a systematic review and meta analysis of safety, satisfaction, and efficiency outcomes. Gastrointestinal Endoscopy. 2020 May 1; 91(5):1015 26.
- Sneyd JR, Absalom AR, Barends CR, Jones JB. Hypotension during propofol sedation for colonoscopy: a retrospective exploratory analysis and meta analysis. British journal of anaesthesia. 2022 Apr 1;128(4):610 22
- Kinugasa H, Higashi R, Miyahar a K, Moritou Y, Hirao K, Ogawa T, Kunihiro M, Nakagawa M. Dexmedetomidine for conscious sedation with colorectal endoscopic submucosal dissection: a prospective double blind randomized controlled

- study. Clinical and Translational Gastroenterology. 2018 Jul 1;9(7):e167.
- 11. Doğanay G, Ekmekçi P, Kazbek BK, Yılmaz H, Erkan G, Tüzüner F. Effects of alfentanil or fentanyl added to propofol for sedation in colonoscopy on cognitive functions: Randomized controlled trial.
- 12. Abu Baker F, Mari A, Aamarney K, Hakeem AR, Ovadia B, Kopelman Y. Propofol sedation in colonoscopy: from satisfied patients to improved quality indicators. Clinical and experimental gastroenterology. 2019 Feb 26:105
- Kinugasa H, Higashi R, Miyahara K, Moritou Y, Hirao K, Ogawa T, Kunihiro M, Nakagawa M. Dexmedetomidine for conscious sedation with colorectal endoscopic submucosal dissection: a prospective double blind randomized controlled study. Clinical and Translational Gastroenterology. 2018 Jul 1;9(7):e167.
- Blayney MR. Procedural sedation for adult patients: an overview. Continuing Education in Anaesthesia, Critical Care & Pain. 2012 Aug 1;12(4):176 80.
- Kang S, Lu J, Zhou HM. Anesthetic strategy for obese patients during gastroscopy: deep sedation or conscious sedation? A prospective randomized controlled trial. Journal of anesthesia. 2021 Aug;35:555 62.
- Johnson KB. Advantages, disadvantages, and risks of TIVA/TCI. Total Intravenous Anesthesia and Target Controlled Infusions: A Comprehensive Global Anthology. 2017:621 31.
- Lu Y, Hao LX, Chen L, Jin Z, Gong B. Systematic review and meta analysis of patient controlled sedation versus intravenous sedation for colonoscopy. International journal of clinical and experimental medicine. 2015;8(11):19793.
- Lee S. Dexmedetomidine: present and future dire ctions. Korean journal of anesthesiology. 2019 Aug 1;72(4):323 30.
- Li A, Yuen VM, Goulay Dufay S, Kwok PC. Pharmacokinetics and pharmacodynamics of dexmedetomidine. Drug development and industrial pharmacy. 2016 Dec 1;42(12):1917 27.
- Sahinovic MM, Struys M M, Absalom AR. Clinical pharmacokinetics andpharmacodynamics of propofol. Clinical pharmacokinetics. 2018Dec;57(12):1539 58.
- Marik PE. Propofol: therapeutic indications and side effects. Currentpharmaceutical design. 2004 Nov 1;10(29):3639 49.
- Seyam SH, Aboelsuod MA, Ahmed IM, Hassan AE. Sedation for ColonoscopyProcedures Using Dexmedetomidine Versus Propofol Fentanyl Infusions: AProspective Randomized Controlled Trial. Turkish Journal of Anaesthesiologyand Reanimation. 2024 Apr;52(2):60.
- Przemyslaw Jalowiecki Robert Rudner Maciej Gonciarz PiotrKawecki Michal Petelenz Piotr Dziurdzik; Sole Use of Dexmedetomidine HasLimited Utility for Conscious Sedation during OutpatientColonoscopy. Anesthesiology 2005; 103:269 273 doi:
- 24. Ghali A, Mahfouz AK, Ihanamäki T, El Btarny AM. Dexmedetomidine versuspropofol for sedation in patients undergoing vitreoretinal surgery under subTenon's anesthesia. Saudi journal of anaesthesia. 2011 Jan 1;5(1):36
- 25. Kavousi E, Shariefnia HR, Pourfakhr P, Khajavi M, Behser esht A.Dexmedetomidine versus propofol in combination with fentanyl for sedationanalgesia in colonoscopy procedures: a randomized prospective study. MiddleEast Journal of Digestive Diseases. 2021 Oct;13(4):328.