

# Wilms Tumor Detection using Deep Learning Approach

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**Abstract**— Wilms tumor, an interesting however imposing pediatric kidney cancer, requests quick and exact finding for ideal helpful mediations and works on quiet results. In this project, we propose an imaginative system for Wilms tumor identification, utilizing advanced deep learning procedures. Our methodology fixates on fastidiously planned deep learning methods prepared on a different dataset of clinical imaging checks including both Wilms tumor cases and typical kidney structures. The deep learning techniques are prepared to independently separate many-sided designs and inconspicuous highlights from clinical pictures, empowering exact recognizable proof of unusual development demonstrative of Wilms tumor. Through thorough preparation, the model is enhanced for increased responsiveness and particularity, guaranteeing strong segregation among dangerous and non-harmful examples. Exhaustive approval on an autonomous dataset assesses the model's generalizability and certifiable dependability. Broad execution assessments against regular symptomatic techniques highlight the prevalence of our deep learning model concerning exactness, awareness, and particularity. This project presents a deep learning-based system for early Wilms tumor discovery, offering a promising road for upgraded demonstrative precision and ideal intercessions. The joining of trend-setting innovation into routine clinical practices can help pediatric oncology, giving a more productive and exact symptomatic worldview that essentially influences patient results.

**Keywords**— Wilms Tumor, Deep learning, Pediatric cancer

## I. INTRODUCTION

Wilms Tumor, often called Nephroblastoma, is a cancer that is seen mostly in kids. It was first observed by a German surgeon Max Wilms in 1899. It is the most common and solid malignant neoplasms in children which occurs in about 90% of the pediatric renal cancers. Quick diagnosis of Wilms tumor is important for effective treatment and better patient results. Traditional methods for Wilms tumor detection have relied on manual examination and interpretation of medical imaging, which can be time-consuming and subjective. Recent advancements in Deep learning approaches have demonstrated considerable potential for boosting accuracy and efficiency of Wilms tumor detection. Nonetheless, endurance rates have dropped in instances of repeat.

Histopathological grouping is vital for treatment delineation and treatment decision. Artificial intelligence could further develop risk delineation by giving more precise and reproducible measurement. AI, especially deep learning (DL)

and neural networks (NNs), has shown guarantee in medical services applications, especially in radiology and urology [1]. A review distinguished remarkable biomarkers for Wilms cancers utilizing Objective and GEO RNA-seq datasets and clinical data. These biomarkers are then examined using machine learning techniques to anticipate, guess and decide their practicality for immunotherapy and chemotherapy [2].

Radiotherapy and chemotherapy can further develop survival rates treating advanced Wilms tumors, although they can grow the secondary malignant cancers. Chemotherapeutic specialists like actinomycin, adriamycin, and vincristine can cause poisonousness, impede hearing and cardiovascular capability, and prompt fringe neuropathy. [2]. However a study reveals that nephroblastoma answers well to neoadjuvant chemotherapy, and deluding conclusion can prompt unseemly chemotherapy regimens. AI techniques, for example, radiomics and deep learning, stand out for quantitatively evaluating clinical imaging information [3]. Radiomics, a novel imaging method, can help in tumor histological subtype characterization, organizing, and visualization examination, giving clinical choice help and a wide application prospect [4]. As of late, MRI-only work processes have been created utilizing MRI-based synthetic CT (sCT) pictures to compute portion testimony, particularly for anatomicies with troublesome MRI-CT enrollment. This dispenses with efficient vulnerabilities and improves on treatment arrangement. sCT has been utilized for different organ transfers, including liver cancer therapy [5]. Cancer stages and histological subtypes are key for therapy results and endurance. Lower stages foresee a positive guess. Regardless of age, tumor weight, and histological type assuming a part. Stage I WT patients might possibly go through nephron-saving resection or nephrectomy without postoperative chemotherapy [4].

In any case, concentrates on sCT age strategies for pediatric patients are scant. Radiologists explore the possibility of an MRI-based treatment arranging work process for youngsters with abdominal cancers utilizing a start to deep learning approach [5].

## II. LITERATURE REVIEW

Wilms tumor is a formidable pediatric renal cancer. The early diagnosis helps in reduction of the relapse rate as it is cured earlier. This helps in reducing the mortality rate when the patients grow up to be teenagers. We have found that there are several studies in the diagnosis of Wilms tumor which helped us in our project. This literature review throws a light on few of the important works related to the Wilms tumor diagnosis and detection during treatment.

Ananda van der Kamp et al.[1] said that WT is more common and frequent children cancer in children and has extremely remarkable pathology, prompting variety of grouping in their study. Man-made reasoning based programmed acknowledgment holds the commitment that this might be finished in a more predictable manner than human eyewitnesses can. They focused on digitized microscopic slides of 72 WT patients, stained with conventional hematoxylin and eosin, made use of neural network to recognize 15 unique normal and growing sections. They illustrate how a deep-learning architecture does this work with greater accuracy, seen by a dice score for the 15 pieces as 0.85, and how this method allow for further automation of Wilms Tumor recognition.

Hanxiang Liu et al.[2] performed a study, that recognizes interesting biomarkers for Wilms growth utilizing Objective and Quality Articulation Omnibus RNA-seq datasets and clinical data. The gamble attributes of these biomarkers were dissected to anticipate the visualization of Wilms patients as well as find their suitability for biological therapy and chemotherapy. The rna-seq transcript genome information, methylation information, and clinical data were retrieved from the Objective data collection. while microarray information from GEO and TCGA skilnet malignant growth RibonucleicAcid sequencing information are downloaded from the UCSC Xena data set. A univariate Cox relative peril relapse model was laid out for each differentially communicated quality (DEG) utilizing endurance information. It was found that, the endurance pace The number of patients with okay grades was much greater than those in the high-risk bunch. The concentrate additionally dissected the prescient connection between Risk score based on both single and multiple components, age, sex, stage, and histological type. The differences in gender, stage, and chance score were quite large, suggesting that they are inverse predictive features for Wilms growth patients. The ROC curve revealed that the two-, three-, and five-year prognostic forecasts would be advised to order productivity than those of sex, age, stage, and histology.

Ilker Ozgur Koska et al. [3] performed a review that planned to foster a powerful grouping model involving children's stomach CT, radiomics and clinical information are utilized to assess neuroblastoma (NB) or Wilms improvement (WT) earlier to surgery. The review included 147 patients, including 90 WT and 57 NB, who were inspected from January 2005 to December 2021. The computer tomography-based office radiologic-radiomics consolidated The model's F1 score was 0.94, accuracy was 0.93, and AUC was 0.96. The model precisely anticipated two patients, which

radiologists couldn't precisely foresee. They mentioned that the model could act as an effortless preoperative sign of Neuroblastoma and Wilms Tumor separation in computer tomography, that ought to be additionally supported in huge scope models, and that artificial intelligence and machine learning approaches may be effectively used to a variety of clinical imaging diagnoses.

Xiao Hui Ma et al.[4] performed a review investigation Includes 118 individuals with wilms tumor which processed contrast-improved CT. examines somewhere in the period of 2014-21 was directed. The patients were partitioned into two gatherings: There are two types of sickness: stage I and nonstage I. Total 1,781 scan properties from 7 component classes were taken from pre-operative intra-venous stage pictures of stomach computer tomography. The Engineered To cope with unbalanced datasets, the Minority Over-Examining Procedure (Destroyed) was used, following by a T-test and the least outright shrinkage. Choice Administrator (Rope) regularization for include determination. A Help Vector Machine (SVM) was conveyed to foster a foreseeing model in light of the chose enlightening elements. The model showed better presentation in the validation dataset, with cross-approved AUC of a 0.79 on the preparation dataset and 0.81 about the test dataset. The AI model precisely anticipated wilms tumor stage-I and non-stage-I illness in child patients prior operation, providing quick as well as easy approach to explore WT phases.

Mateusz C. Florkow et al.[5] performed a study of 66 pediatric patients with WT or NB(age:  $4 \pm 2$  years) utilized X-ray and computer tomography procurement to evaluate the possibility of attractive reverberation imaging (X-ray)- just therapy making arrangements for photon and proton radiation in kids with stomach growths. X-ray forces were switched over completely to CT Hounsfield units (HU) utilizing an UNet-like brain organization. The review assessed CT-to-sCT picture likeness utilizing mean blunder, mean outright mistake, top sign to-clamor proportion, and Dice comparability coefficient. Manufactured CT dosimetric exactness was confirmed against computer tomography dosimetric accuracy was proven against computer tomography-based portion dispersions for volumetric-regulated circular segment treatment (VMAT) and power balanced pencil-shaft examining (PBS). They found that most dosimetric contrasts were inside clinically satisfactory models for photon and proton radiotherapy, showing the achievability of a X-ray just work process for pediatric patients with stomach cancers. The profound gaining based model produced exact sCT from arranging T1w-and T2w-MR pictures, showing the achievability of a X-ray just work process for pediatric patients with stomach cancers.

Elwira Szychot et al.[6] mentioned in their study that Wilms' tumor, influencing one of every 10,000 youngsters, is a typical renal disease. It has a high long haul endurance rate, with 90% for confined sickness and 75% for metastatic infection. Effective administration requires cautious organizing and coordinated effort between oncologists, specialists, radiologists, pathologists, and radiation oncologists. Current treatment conventions use risk task to

limit harmfulness and further develop results for high-risk patients. Future endeavors will focus on biomarkers and and further develop results for patients with unfavourable histology and repetitive sickness.

Mujie Li et al.[7] performed the review, that planned to create and approve an estimate the possibility of myelosuppression caused by chemotherapy (CIM) in kids with Wilms Tumor(WT) prior treatment is managed. The model depended on information from 1433 chemotherapy rounds were administered to 437 children with WT. Six ML calculations were utilized to build the models, and the prescient adequacy of these models was assessed. It was found that in 58.5% of treatment cycles, there was grade  $\geq 2$  CIM. The outrageous slope helping (XGB) model was distinguished as the best model because of its high prescient proficiency and security. EGG whites, basic phosphatase, hemoglobin, white platelet count, and coadministration of extremely toxic chemotherapeutic medicines were the five elements that contributed most to the model. XGB model was created interestingly and has great prescient execution and dependability. They mentioned that model's true capacity for clinical applications could prompt the augmentation of CIM expectation models to other pediatric malignancies.

Israa Sharaby et al.[8] performed a review that meant to foster a preoperative chemotherapy helped expectation framework for preoperative chemotherapy reaction in Wilms' cancers, a typical renal growth in youngsters. The framework included 63 patients matured 6-14 years, who were investigated utilizing contrast-improved registered tomography imaging. The framework extricated surface, shape, and usefulness based highlights from the cancers before chemotherapy. The framework comprises of six stages: portraying the growths' pictures, portraying the surface utilizing textural highlights, separating shape highlights utilizing parametric circular music model, catching power changes to depict the cancers' usefulness, applying highlights combination in view of removed includes, and determining whether or not the most recent forecast was responsive utilizing a customized help vector machine classifier. Framework accomplished a general exactness of 95.24%, with 95.65% awareness and 94.12% particularity. This research shows that Wilms tumors can be anticipated early with preoperative chemotherapy using new imaging indicators and a predictive order model, which could result in customized treatment regimens.

Sabine Müller et al.[9] gave the overview of the usage of The MRI scans of the patients are evaluated using algorithms for DL and ML. The prediction of Wilms tumor before and after the chemotherapy administration greatly varies from the ground truth, as much as 0.15 difference in dice score when observed by human experts. Since the tumor size cannot be assessed properly after the chemotherapy has been administered due to the fact, that the tumors' outlines and density cannot be captured by human eye properly from MRI scans alone. However with the usage of machine learning algorithms the dice score is more than that of human experts, and the difference in the dice score before and after the chemotherapy is also less compared to the evaluation of

human experts. The dice score of Human experts before and after chemotherapy are  $0.93 \pm 0.05$  and 0.85 respectively. Meanwhile the maximum dice score of the machine learning algorithms before and after chemotherapy are 0.92 and 0.81 respectively. Hence, it can be said that usage of refined deep learning techniques can bring out accurate results in medical diagnosis.

Yaxin Deng et al.[10] performed a review to evaluate the effectiveness of CT radiomics in distinguishing pediatric patients' CCSK from WT. The review comprised 83 cases of wilms tumor and 33 cases of clear cell sarcoma of kidney, defined a preparation as well as test set. Imaging highlights from the nephrographic stage were examined, including growth measurement, the larger peritumoral blisters and the ratio of CTmax to CT renal veins. To develop a strategic relapse model, radiomics highlights from the corticomedullary stage were utilized. Exactness, 95 percent confidence span (CI) and region under the bend (AUC) were used to evaluate the model's presentation. Model areas of strength for showed in the preparation set with an AUC of 0.889 and an exactness of 0.864. After fivefold cross-approval, the AUC stayed high at 0.863 and a precision 0.852. The model produced an accuracy of 0.857 and an AUC of 0.792 in the test set. CT radiomics ended up being analytically significant for recognizing WT and CCSK in kids.

### III. MATERIALS AND METHODS

#### A. Dataset

This project uses the custom made dataset consisting of the MRI scans of the Wilms tumor patients and patients suffering from other pediatric tumors. The images of the MRI scans were taken from Radiopedia.org a non profit organization where the reports, cases and diagnosis of the radiomics are discussed with the expert opinions. We have considered total of 20 patients data which consisted of Axial c portal venous phase MRI scans consisting of 60 images intotal which were later augmented to expand the dataset consisting of near 1000 images. We have used this data for research and model development purpose only. The dataset is parted in testing to validation ratio of 8:2.

#### B. Software and Tools

- Python: The programming language used for creating and executing the profound learning models.
- TensorFlow and Keras: These deep learning systems are utilized for model turn of events, preparing, and assessment.
- OpenCV: A library utilized for different picture handling undertakings, for example, resizing, increase, and post-handling.
- PIL: It is used to convert the image extensions to jpeg.
- Google collab: a code execution notebook is utilized to run and execute the created code with the model and dataset.

### C. Preprocessing

To guarantee consistency and upgrade model execution, the MRI scans checks go through a few preprocessing steps:

- Normalization: Force upsides of the MRI scans checks are standardized to a scope of 0 to 1, normalizing the information.
- Resizing: Pictures are resized to 416x416 pixels to match the info prerequisites of the Consequences be damned model.
- Augmentation: Information increase methods like revolution, flipping, and zooming are applied to the preparation dataset. These techniques increase the changeability of the dataset, assisting with forestalling overfitting and work on the model's capacity to sum up to inconspicuous information.

### D. Model Architecture

The essential model utilized for Wilms tumor location is the You Only Look Once (YOLO) architecture, explicitly YOLOv8, which is famous for its productivity and exactness in object identification errands.

### E. Labeling

Bounding boxes are explained for tumor boundaries in the MRI scans checks. Each output is related with at least one jumping boxes, demonstrating the presence and exact area of Wilms tumors.

### F. Evaluation Metrics

Model execution is thoroughly assessed utilizing a few measurements:

- Accuracy - the ratio of correctly predicted instances (both Wilms tumor and non-Wilms tumor) to the total no. of instances in the dataset.

$$\text{Accuracy} = \frac{\text{no.of correct predictions}}{\text{total no.of predictions}}$$

- Precision - determines the proportion of all the model's positive predictions that are real positive findings (identifies Wilms tumors accurately).

$$\text{Precision} = \frac{\text{true positives}}{(\text{true positives} + \text{false positives})}$$

- Recall / sensitivity - it measures the proportion of actual positive instances (Wilms tumors) that are correctly predicted by the model.

$$\text{Recall} = \frac{\text{true positives}}{(\text{true positives} + \text{false negatives})}$$

- F1 score - It's particularly useful when you need to consider both precision and recall simultaneously

$$\text{F1 score} = 2 \times \frac{\text{Precision} \times \text{recall}}{(\text{Precision} + \text{recall})}$$

- Dice coefficient - Higher numbers indicate better overlap; the range is 0 to 1.

$$\text{DC} = \frac{(2 \times \text{Area of overlap})}{(\text{Area of predicted region} + \text{Area of ground truth region})}$$

### G. Validation and Testing

- Cross-Validation: A 5-overlap cross-validation is performed on the preparation set to guarantee model power and relieve the risk of overfitting.
- Independent Test Set: The model's generalizability is surveyed utilizing an autonomous test set, giving a fair assessment of execution.

### H. Post-Processing

Non-Greatest Concealment (NMS) is applied to the anticipated bouncing boxes to wipe out copy location and hold the most certain forecasts. Moreover, morphological tasks are utilized to refine the edges of the bounding boxes, working on the precision of tumor confinement.

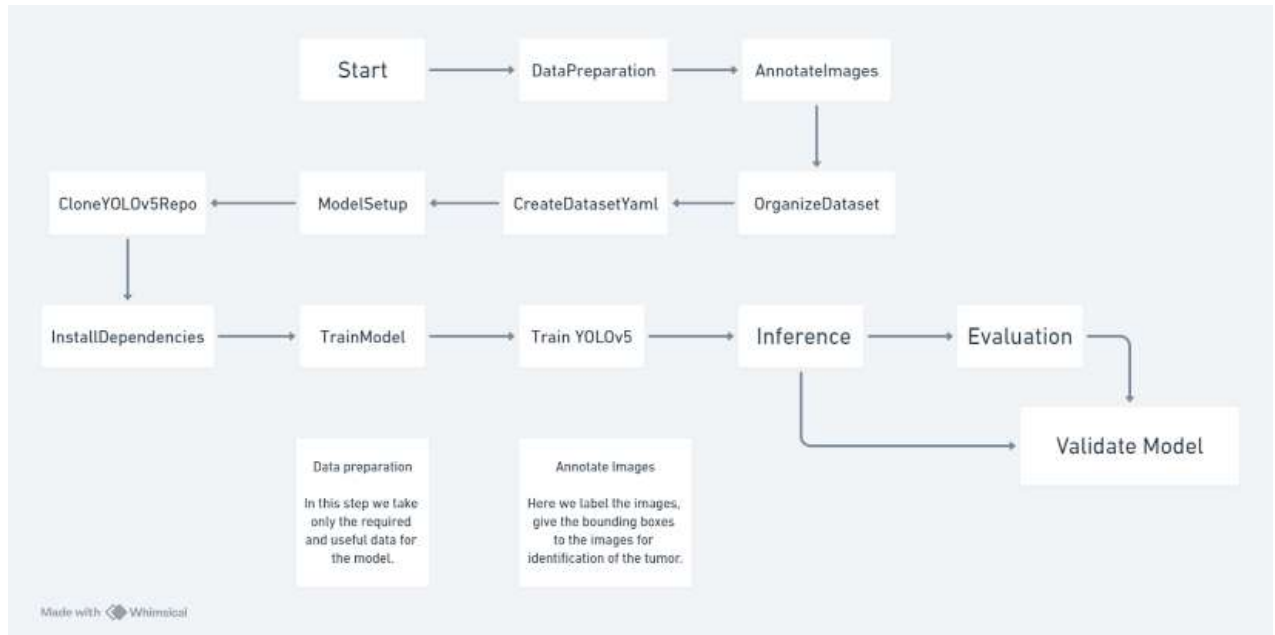
## IV. PROPOSED METHOD

The proposed method uses You Only Look Once (YOLO) deep learning architecture to detect the presence of the Wilms tumor in the MRI scans. This method uses the custom dataset from the images of the MRI scans were taken from Radiopedia.org a non profit organization where the reports, cases and diagnosis of the radiomics are discussed with the expert opinions. We have considered total of 20 patients data which consisted of Axial c portal venous phase MRI scans consisting of 60 images in total which were later augmented to expand the dataset consisting of near 1000 images.

The YOLOv8 design is picked for its pace among. speed and exactness, making it appropriate for continuous article identification assignments. You Only Look Once, is a creative deep-learning model that reformulates object location as a solitary relapse issue, An assessment of full pictures that predicts the bounding boxes and course probabilities in a clear way. This approach diverges from conventional strategies that utilization a locale proposition organization, fundamentally decreasing calculation time and making YOLO highly proficient.

### A. Key Components of YOLOv4

- Backbone - CSPDarknet53: The backbone network, CSPDarknet53, fills in as the element extractor, giving high-goal include maps significant for recognizing little and complex designs like cancers. This network is planned with Cross-Stage Partisl (CSP) associations, upgrading angle stream and further developing the learning capacity of the network without altogether expanding computational intricacy.



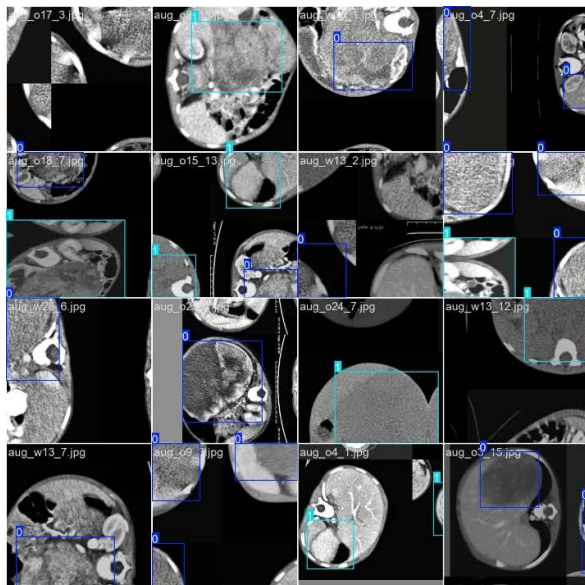
**Fig 1. Implementation**

### B. Advantages of YOLO

- **Neck - PANet and SPP Layers:** The neck of the YOLOv8 model integrates PANet and Spatial Pyramid Pooling layers. PANet improves the data stream between various layers by considering better component combinations, and upgrading the model's capacity to distinguish objects at different scales. SPP layers, then again, help in keeping up with spatial data, permitting the model to really deal with varieties in tumor size and shape.
- **Head:** The location head is liable for anticipating bouncing boxes, class probabilities, and objectness scores. It yields three sizes of forecasts, obliging tumors of different sizes and guaranteeing exact limitation and grouping.

The following key points are the advantages of the YOLO architecture-

- **Speed:** YOLO is especially quick, making it appropriate for continuous applications. This is especially valuable in clinical settings where convenient navigation is basic.
- **Unified Architecture:** YOLOs start to finish preparing and expectation pipeline improves on the recognition cycle, diminishing the requirement for complex post-handling steps.
- **Accuracy:** Regardless of its speed, Consequences be damned keeps up with high precision by really adjusting accuracy and review, making it solid for clinical picture examination.



**Fig 2. Training image**

### C. Training Process

#### 1) Hyperparameters:

- Optimizer:** The Adam streamlining agent is utilized with an underlying learning pace of 0.001. Adam's versatile learning rate capacities make it appropriate for training deep neural networks.
- Loss Function:** The YOLO loss function is a blend of classification loss, confinement loss(bounding box relapse), and confidence loss (objectness score). This composite loss guarantees adjusted enhancement, zeroing in on exact bounding box relapse and sure tumor identification.
- Epochs and Batch Size:** The model is prepared for 50 epochs, with early halting in light of validation loss to forestall overfitting. A group size of 16 is utilized, adjusting computational productivity and intermingling dependability.

## V. RESULTS AND DISCUSSIONS

This model was applied to 1000 MRI scan images taken from a custom-built dataset augmented from the data of the 20 patients from radiopedia.org. We have used labeled data for augmentation and obtained the related augmented data. This augmented data can help predict the unknown and different cases of wilms tumor from other tumors. While we evaluated the model we found that it has 97.40% accuracy, this is the big step towards more accurate diagnosis. We have also found that the precision of the model is 96.55 % which needs to be worked on. The recall value remained 98.26% while F1 score and Dice coefficient are 0.97 and 0.97 respectively.

Metric	Value
Accuracy	97.40%
Precision	96.55%
Recal value	98.26%
F1 score	0.97
Dice coefficient	0.97

Table 1. Results

We can also visualize the training loss and validation loss from the graph below. The training loss decreased as the number of epochs increased.

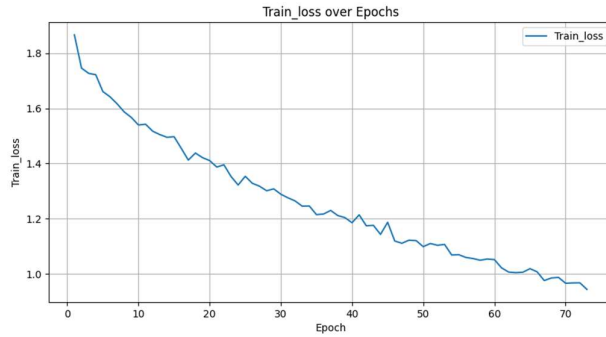


Fig 3. Training Loss

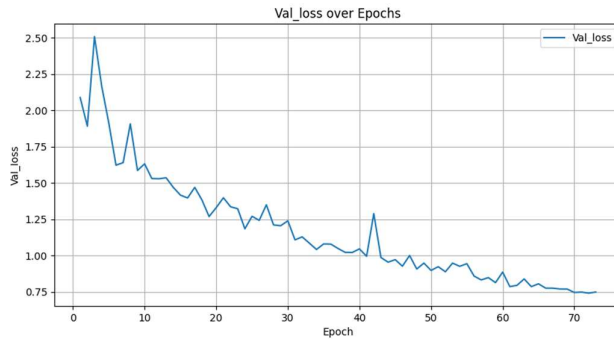


Fig 4. Validation Loss

The dataset can be more organised and well processed to yield better results with this model. Also many other parameters can also be included in order to improve the Wilms tumor diagnosis' precision, usage of pyradiomics will also help to extract many features out of the MRI images, this also helps in increasing metrics of the developed model.

## VI. CONCLUSION

The proposed technique for Wilms tumor identification in MRI scans utilizes the YOLOv8 algorithm, planning to convey a fast, exact, and computerized answer for clinical applications. YOLOv8's architecture, which incorporates CSPDarknet53 as the backbone, alongside PANet and SPP layers in the neck, guarantees high-goal highlight extraction and powerful multi-scale highlight combination. This development permits the model to identify tumors of shifting sizes with high accuracy and review effectively.

The technique's utilization of exhaustive preprocessing steps, including normalization, resizing, and augmentation, improves the model's vigor and generalizability. Preparing the model with the Adam enhancer and an even misfortune capability further guarantees precise bounding box relapse and accurate tumor location.

YOLO's start to finish engineering offers huge benefits in speed and effectiveness, making it especially reasonable for ongoing applications in clinical settings. The model's fast and exact growth limitation abilities can incredibly help radiologists, upgrading analytic precision and working on persistent results through opportune and precise intercessions.

This YOLOv8-based approach for Wilms Tumor discovery in MRI scans presents a promising tool for clinical imaging. Its equilibrium of speed, exactness, and productivity features its capability to altogether improve the proper diagnosis of the tumor and treatment plan, eventually adding to better medical services results.

## REFERENCES

- [1] van der Kamp A, de Bel T, van Alst L, Rutgers J, van den Heuvel-Eibrink MM, Mavinkurve-Groothuis AMC, van der Laak J, de Krijger RR. Automated Deep Learning-Based Classification of Wilms Tumor Histopathology. *Cancers*. 2023; 15(9):2656. <https://doi.org/10.3390/cancers15092656>
- [2] Hanxiang Liu, Chaozhi Tang, Yi Yang, "Identification of Nephrogenic Therapeutic Biomarkers of Wilms Tumor Using Machine Learning", *Journal of Oncology*, vol. 2021, Article ID 6471169, 14 pages, 2021. <https://doi.org/10.1155/2021/6471169>
- [3] Koska, I.O., Ozcan, H.N., Tan, A.A. *et al.* Radiomics in differential diagnosis of Wilms tumor and neuroblastoma with adrenal location in children. *Eur Radiol* (2024). <https://doi.org/10.1007/s00330-024-10589-8>
- [4] Ma X-H, Shu L, Jia X, Zhou H-C, Liu T-T, Liang J-W, Ding Y-s, He M and Shu Q (2022) Machine Learning-Based CT Radiomics Method for Identifying the Stage of Wilms Tumor in Children. *Front. Pediatr.* 10:873035. doi: 10.3389/fped.2022.873035
- [5] Mateusz C. Florkow, Filipa Guerreiro, Frank Zijlstra, Enrica Seravalli, Geert O. Janssens, John H. Maduro, Antje C. Knopf, René M. Castelein, Marijn van Stralen, Bas W. Raaymakers, Peter R. Seevinck, Deep learning-enabled MRI-only photon and proton therapy treatment planning for paediatric abdominal tumours, *Radiotherapy and Oncology*, Volume 153, 2020,



- Pages 220-227, ISSN 0167-8140,  
<https://doi.org/10.1016/j.radonc.2020.09.056>.
- [6] Szychot, E., Apps, J., & Pritchard-Jones, K. (2014). Wilms' tumor: biology, diagnosis and treatment. *Translational pediatrics*, 3(1), 12–24. <https://doi.org/10.3978/j.issn.2224-4336.2014.01.09>
  - [7] Li M, Wang Q, Lu P, Zhang D, Hua Y, Liu F, Liu X, Lin T, Wei G, He D. Development of a Machine Learning-Based Prediction Model for Chemotherapy-Induced Myelosuppression in Children with Wilms' Tumor. *Cancers*. 2023; 15(4):1078. <https://doi.org/10.3390/cancers15041078>
  - [8] Sharaby I, Alksas A, Nashat A, Balaha HM, Shehata M, Gayhart M, Mahmoud A, Ghazal M, Khalil A, Abouelkheir RT, et al. Prediction of Wilms' Tumor Susceptibility to Preoperative Chemotherapy Using a Novel Computer-Aided Prediction System. *Diagnostics*. 2023; 13(3):486. <https://doi.org/10.3390/diagnostics13030486>
  - [9] Sabine Müller, Iva Farag, Joachim Weickert, Yvonne Braun, André Lollert, Jonas Dobberstein, Andreas Hötter, and Norbert Graf "Benchmarking Wilms' tumor in multisequence MRI data: why does current clinical practice fail? Which popular segmentation algorithms perform well?," *Journal of Medical Imaging* 6(3), 034001 (19 July 2019). <https://doi.org/10.1117/1.JMI.6.3.034001>
  - [10] Deng, Y., Wang, H. & He, L. CT radiomics to differentiate between Wilms tumor and clear cell sarcoma of the kidney in children. *BMC Med Imaging* 24, 13 (2024). <https://doi.org/10.1186/s12880-023-01184-2>
  - [11] "Wilms tumor—childhood: Statistics", 2022, [online] Available: <https://www.cancer.net/cancer-types/wilms-tumor-childhood/statistics>.
  - [12] E. Szychot et al., "Wilms' tumor: biology diagnosis and treatment", *Translational Pediatrics*, vol. 3, no. 1, pp. 12, 2014.
  - [13] The SIOP Renal Tumour Study Group Paediatric renal tumours: Perspectives from the SIOP-RTSG. *Nat. Rev. Urol.* 2017;14:3–4. doi: 10.1038/nrurol.2016.247.
  - [14] Chahal A., Gulia P. Machine learning and deep learning. *Int. J. Innov. Technol. Explor. Eng.* 2019;8:4910–4914. doi: 10.35940/ijitee.L3550.1081219.
  - [15] Esteva A., Kuprel B., Novoa R.A., Ko J., Swetter S.M., Blau H.M., Thrun S. Dermatologist-level classification of skin cancer with deep neural networks. *Nature*. 2017;542:115–118. doi: 10.1038/nature21056.
  - [16] Yadav A.S., Kumar S., Karetla G.R., Cotrina-Aliaga J.C., Arias-González J.L., Kumar V., Srivastava S., Gupta R., Ibrahim S., Paul R., et al. A Feature Extraction Using Probabilistic Neural Network and BTFSC-Net Model with Deep Learning for Brain Tumor Classification. *J. Imaging*. 2022;9:10. doi: 10.3390/jimaging9010010.
  - [17] Kingma, D. P., & Ba, J. (2014). Adam: A method for stochastic optimization. *arXiv preprint arXiv:1412.6980*.
  - [18] Popov, S. D., Sebire, N. J., & Vujanic, G. M. (2016). Wilms' tumour—histology and differential diagnosis.
  - [19] Ronneberger, O., Fischer, P., & Brox, T. (2015). U-net: Convolutional networks for biomedical image segmentation. In Medical image computing and computer-assisted intervention—MICCAI 2015: 18th international conference, Munich, Germany, October 5-9, 2015, proceedings, part III 18 (pp. 234-241). Springer International Publishing.
  - [20] Ehrlich, P., Chi, Y. Y., Chintagumpala, M. M., Hoffer, F. A., Perlman, E. J., Kalapurakal, J. A., Warwick, A., Shamberger, R. C., Khanna, G., Hamilton, T. E., Gow, K. W., Paulino, A. C., Grattias, E. J., Mullen, E. A., Geller, J. I., Grundy, P. E., Fernandez, C. V., Ritchey, M. L., & Dome, J. S. (2021). Combination Chemotherapy and Surgery in Treating Young Patients With Wilms Tumor (AREN0534) [Data set]. The Cancer Imaging Archive. <https://doi.org/10.7937/TCIA.5M9S-6Y97>
  - [21] B. K. Sethi, D. Singh, S. K. Rout and S. K. Panda, "Long Short-Term Memory-Deep Belief Network-Based Gene Expression Data Analysis for Prostate Cancer Detection and Classification," in *IEEE Access*, vol. 12, pp. 1508-1524, 2024, doi: 10.1109/ACCESS.2023.3346925.
  - [22] N. Routray, S. K. Rout, B. Sahu, S. K. Panda and D. Godavarthi, "Ensemble Learning With Symbiotic Organism Search Optimization Algorithm for Breast Cancer Classification and Risk Identification of Other Organs on Histopathological Images," in *IEEE Access*, vol. 11, pp. 110544-110557, 2023, doi: 10.1109/ACCESS.2023.3322222.
  - [23] Sahu, B., Mohanty, S. and Rout, S., 2019. A hybrid approach for breast cancer classification and diagnosis. *EAI Endorsed Transactions on Scalable Information Systems*, 6(20), DOI:10.4108/eai.19-12-2018.156086
  - [24] Rout, S.K., Kumar, K.S., Sahu, R., Barik, S. and Pradhan, S., 2024, March. Tropical Cyclone Detection and Tracking Using YOLOv8 Algorithm. In 2024 1st International Conference on Cognitive, Green and Ubiquitous Computing (IC-CGU) (pp. 1-6). IEEE
  - [25] K. Sulthana, F. Begum, G. B. Regulwar, S. K. Rout, V. Dasari and A. B, "Parkinson's Disease Prediction using XGBoost and SVM," *IEEE Xplore Digital Library* 979-8-3503-4488-2 @2023 2nd International Conference on Ambient Intelligence in Health Care (ICAHC), Bhubaneswar, India, 2023, pp. 1-6, doi: 10.1109/ICAHC59020.2023.10431437.
  - [26] G. B. Regulwar, N. Gudipati, M. Naik, N. Kathirisetty, L. Madiga and E. R. Kumar, "Audio to Sign Language Translator," 2023 2nd International Conference on Ambient Intelligence in Health Care (ICAHC), Bhubaneswar, India, 2023, pp. 1-6, doi: 10.1109/ICAHC59020.2023.10431466