

# Forecasting the risk factors of COVID-19 through AI feature fitting learning process (FfitL-CoV19)

**Abstract:** In the present day situation, decreasing COVID-19 risk elements also can moreover need to probably check the abilities to provide and test AI-primarily based total models for COVID-19 severity prediction. This work aimed to select the maximum affecting abilities of COVID-19 chance elements and enhance the functionality of the AI method for COVID-19 chance elements primarily based totally on the chosen abilities. In this paper we've proposed a way to find out whether or not an affected person has a chance of COVID-19 through the usage of an AI characteristic becomes into the feature-fitting learning process (FfitL-CoV19), thinking about a couple of symptoms. In the proposed AI characteristic becoming into the learning process, textual information has been classified into classical and ensemble system learning algorithms. Feature engineering has become completed the use of the AI technique and functions skills have been provided to conventional and ensemble system learning classifiers. The proposed hybrid technique significantly improves preceding solutions and achieves an advanced preferred standard overall performance of efficiency. In the future, the proposed technique may be used to design a structural model as an outline for the multiplicative nature of COV-19 infections.

**Keywords:** AI, COV-19, ML, characteristic fitting, risk factor, feature, ensemble, learning, infection, hybrid, affect, people.

## I. INTRODUCTION:

The pandemic has influenced health problems and maximum human beings' lives—socially and economically [1] [2]. Radiologists are learning to identify COV-19 instances based on chest CT scan image exams, as it is manual and time-consuming, because of the widespread COV-19 and having to examine a lot of cases in less time. This widespread has increased the infection rate at high-risk levels, causing scientists to make it unable to find it manually. There is a need for the development of learning techniques that can analyze and identify the presence of COV-19 severity and can able to distinguish between different COV-19-related disease presences, but it is difficult to have a better and faster method to analyze them at that moment.

There is a lot of use for chest radiology evaluations, and it's a cheap way for digital diagnostics. There is a dearth of high-quality, publicly available, intelligent datasets. The use of pulmonary X-rays in a wide variety of medical settings presents substantial diagnostic and identification challenges that need the use of therapeutic aids. For very large collections of x-ray pictures, the dearth of identifying features makes interpretation difficult. While the prevalent use of lung diagnostic imaging techniques [19] in contemporary medical amenities has improved adaptability and sped up medical imaging evaluations, it has also boosted the economic development of practitioners with specialized skills to characterize the x-ray images in line with the particular patterns reported.

In this study, an observation of the AI method for COV-19 assessment through the utilization of chest X-ray and CT images are identified. To improve the learning method, the researchers have to analyze all the symptoms of contagions related to COV-19 infections by predicting its varying parameters globally and scientifically and should be able to provide better decision-making standards through a defined in a better way effective analysis action on identified symptoms. These can be accomplished through improved scientifically proven mathematical models that have to be developed.

The facts include clinical opinions within the shape of textual content in this paper; we're classifying that visual information into a group of illnesses regions such that it could assist in detecting coronavirus from in advance clinical symptoms. Artificial Intelligence (AI) is becoming a better learning model to research COV-19 symptoms, which has become supporting research to radiologists and scientists, providing a better improvement in healthcare scan data analysis and providing improved patient monitoring care methods. AI plays a significant role in providing the best solution for COV-19 symptoms. Through AI methods, clinical COV-19 data of chest

CT images are analyzed based on the available dataset and scan images, doing machine learning approach and data support vector machine analysis and these can be applied to test the COV-19 risks at potential levels at different levels of critical levels.

By analyzing Computed tomography and pulmonary X-ray pictures from a specific data set and comparing them with specified models, we were able to utilize an ML technique to identify language based on humans with non-infected and Cov-19 affected cases (InceptionV3-and-DensNet).

The remaining paper is categorized into sections: sect 2 provides a previous works. The recommended work is analyzed in sect 3. The implementation analysis are discussed in sect 4. And the conclusions of this work are discussed in sect 5 with the future scope.

## **II. Previous Works:**

Several researchers have proposed automated COV-19 screening through the technique of reading chest radiography or CT images, figuring out the path morphological changes due to Covid-19 within the patient's chest. Other authors, instead, have recommended the evaluation of coughing or respiratory signals, to research the presence of changes because of the COV-19 infection. A few previous types of research, instead, have indicated clarifications for the detection of COV-19 problems primarily based totally on an evaluation of voice samples.

An AI approach is used for the epidemiologic susceptibility infection (SI) model to estimate the spread of COV-19 [3]. The SI approach and its intensification, the SIR model, are both standard outbreak models for modeling and forecasting the diffusion of contagious diseases; in these models, S stands for the form of vulnerable people, I denote for the form of contaminated individuals, and R stipulates the recovered cases.

The consequences of COV-19 dysfunction on human lungs are discussed by the authors in [5]. Next, the authors presented a COV model to extract the classification capabilities of CNN [6].

A DTL (deep transfer learning) of the CNN and RNN Models was developed and assessed [18] to detect and identify COV-19 X-ray chest images. Each improved outcome is achieved using the suggested models. When compared to other models, our model's prediction using CNN, COV-19 is the most accurate and precise. Both the accuracy and precision scores are quite high, at 97.7% and 96.4% respectively. As the gap between the two (training and validation) narrowed, the optimal training procedure emerged. As the F-measure rose to 0.97, a reliable COVID-19 detector was developed. The model performed quite well according to AUC metrics, reaching 0.9. The measure AUC's outcome is shown in Figure 1. As a result, the x-ray data-trained COV-19 diagnostic model delivers higher performance metrics

While ML has only been employed in a small number of COV-19 investigations [8, 9], which has been used to develop a model that compares the characteristics of healthy individuals and affected SARS-CoV-2 victims. The identification of COV-19, its forecasting, and the development of countermeasures [12, 13] are the subject of other studies [10, 11]. In many instances, additional resources are not required for an assessment since all relevant information is already included in the patient's medical record. Using a collection of training, validation, and test instances, [10] develops a machine learning model to predict how an infection's severity will change over time. An SVM set of algorithms is used for prediction, and a genetic set of algorithms (GA) is used in the feature choice stage. In [11], we employed ML to explore the connection between distinct cardiac signs and symptoms and the severity of COV-19 patients [12]. However, some studies show that individuals are less likely to follow preventative recommendations such as isolation [13], good cleanliness [14], and immunization [15].

## **III. Proposed Model:**

Contributions of this work are summarized in the subsequent points:

- (i) Present a dataset built based totally, mostly on different scientific features consisting of the patient's medical history.
- (ii) Feature-fitting into technique based totally, mostly on features function assessment in the detection of COVID-19 virus
- (iii) Propose a risk and infection severity prediction method for COVID-19 patients within the early level of detection.

#### A. AI function fitting into a learning process:

In this section, an ANN-optimization hybrid model for Extraction Of features is suggested. This model combines the advantages of both methods. The predicted approach is evaluated to the pre-trained InceptionV3 and DensNet for making diagnoses on COVID-19 instances.

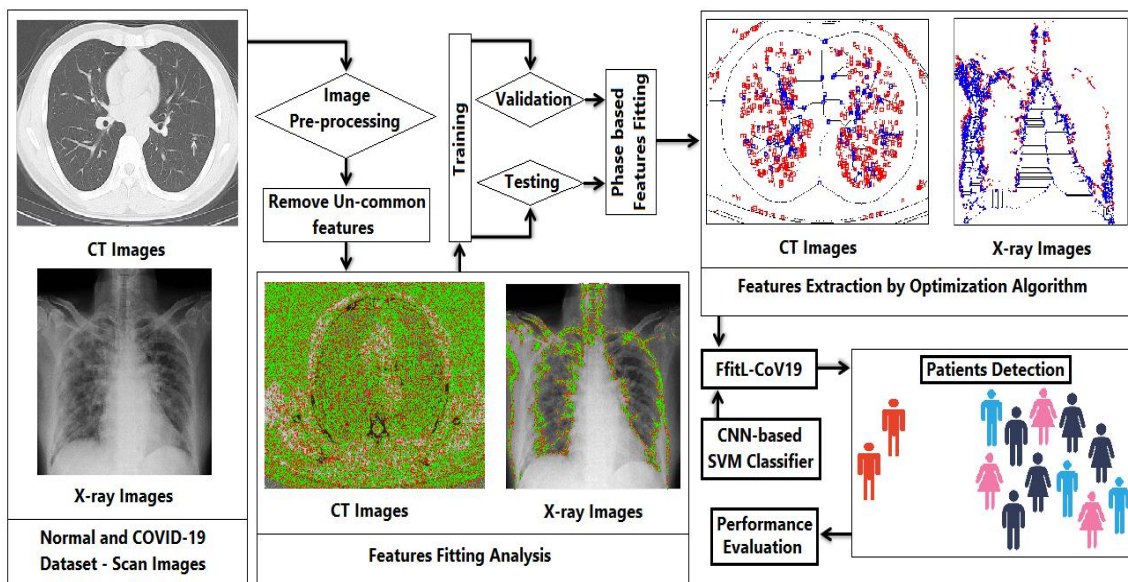


Figure 1: Proposed FfitL-CoV19 process.

#### Algorithm 1: Features Extraction by Optimization Algorithm for FfitL-CoV19 process

Input: Features fitting analysis-CT images ( $CT_i$ ) and X-ray images ( $XR_i$ ), Phase based Features Fitting-trained images ( $T_i$ ), learning rate ( $L_r$ ), population size ( $P_s$ ), number of iterations ( $N_i$ ), and the cumulative number of cases ( $C$ )

Output: Trained FfitL-CoV19 model to classify COVID-19 images

Begin

*Phase-based Features Fitting:*

$\hat{F}$  [ $CT_i$  :  $XR_i$ ]: initialize  $\hat{F}$  as a fitting value

For each image

Do process  $P=SVM(N_i, P_s, L_r)$

for:  $i=1: N_i$

for:  $j=1: P_s$

Update-the-feature-fitting-parameter through  $L_r$

end-for:

end-for:

Find the best  $\hat{F}$ :

for:  $i=1: N_i$

```

for: j=1: Ps
Assign Ti in the range of [0: Ni]
If  $\hat{F} < T_i$ , then
Extract the features with Ti and do-again-Training process
Else If  $\hat{F} > T_i$ , then
Extract the features with  $\hat{F}$  and perform the CNN process
Else  $\hat{F} = T_i$ , then
Optimize the features with the FfitL-CoV19 method
end-if:
end-for:
end-for:
Update-the- $\hat{F}$ -and-the FfitL-CoV19 values
FfitL-CoV19 method:
for: each  $\hat{F}$  feature
Do method M= FfitL( $\hat{F}$ , Ps, Lr, S(t), I(t), R(t), D(t))
Learning rate:  $L(\hat{F}, \tilde{F}) = [(1 - \hat{F}) \log(1 - \tilde{F}) - \sum \hat{F} * \log \tilde{F}]$ 
where  $\tilde{F}$  are the predicted features
Features Fitting parameter:  $Ffit = (P_s * \tilde{F}) + L_r$ 
Classify risk factor:  $\gamma(FfitL) = [C\Delta I(t)^T C\Delta I(t)]^{-1} \min C\Delta I(t)^T C\Delta D(t)$ 
Classify recovery rate:  $\beta(FfitL) = [C\Delta I(t)^T C\Delta I(t)]^{-1} \min C\Delta I(t)^T C\Delta R(t)$ 
The optimal value of infection rate:  $\alpha = [w_i \gamma(FfitL) : w_j \beta(FfitL)]$ 
where  $w_i$  and  $w_j$  are weighing factors of features
Features Detection:  $FfitL = C\Delta I(t)^\alpha C\Delta R(t)^\beta \min C\Delta D(t)^\gamma$ 
Update the FfitL
end
end
end
end

```

#### IV. Results and Discussions:

Table 5 showed how recommended measures of performance in comparison to those of existing works.

##### A. Dataset Evaluation

**Table 1** shows the datasets used and the description of the dataset portioning is shown in **Table 2**.

Table 1: computed tomography scans and X-rays of COV-19 patients and healthy individuals

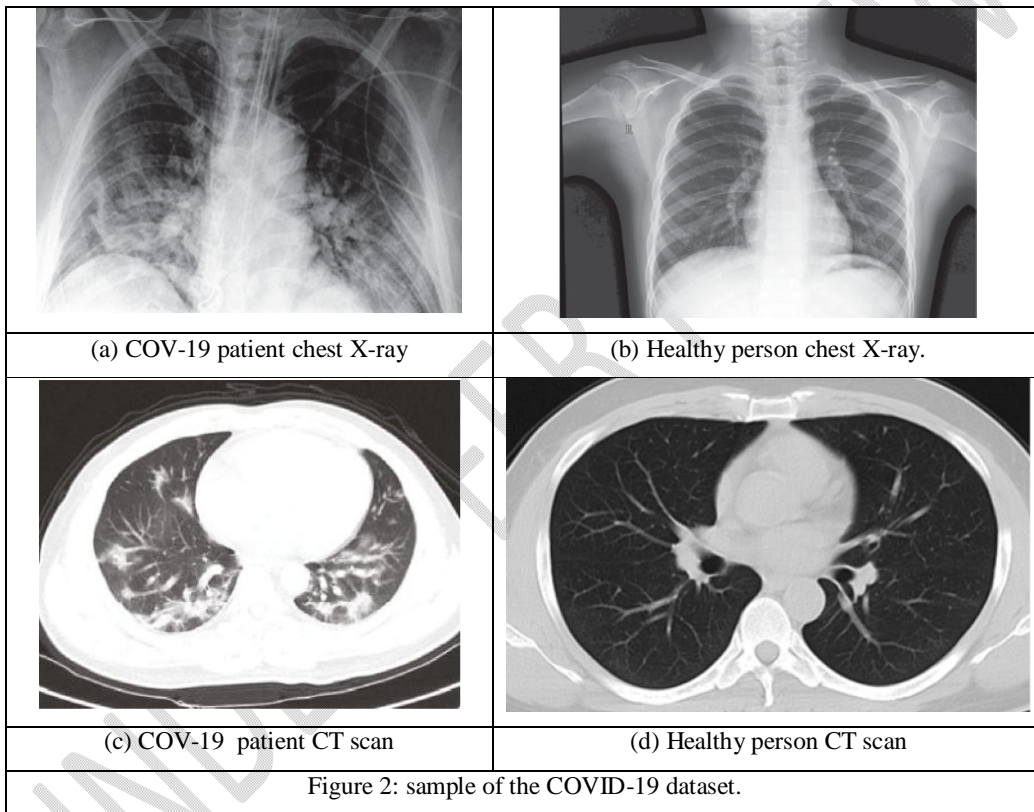
| Type    | Dataset Name                                      | Feature Fitting Data | COV-19 | Non-COV-19 |
|---------|---|----------------------|--------|------------|
|         | COV-19 infected Vs normal cases X-ray images      | Ffit_1               | 1011   | 1308       |
| X-ray   | Auxiliary data X-ray images of COV-19             | Ffit_2               | 1392   | 789        |
|         | COV-19 X-ray images                               | Ffit_3               | 223    | 232        |
|         | COV-19 Vs normal Computed tomography chest images | Ffit_4               | 337    | 365        |
| CT-scan | Computed tomography scan images of SARS-COV-2     | Ffit_5               | 621    | 1166       |
|         | COV-19 Computed tomography scan images            | Ffit_6               | 61     | 80         |

Table 2: Dataset portioning

| Feature Fitting Data | COV-19      |                |              |       | Non-COV-19 |     |     |       |
|----------------------|-------------|----------------|--------------|-------|------------|-----|-----|-------|
|                      | T(Training) | V (Validation) | Te (Testing) | Total | T          | V   | Te  | Total |
| Ffit_1               | 720         | 146            | 145          | 1011  | 912        | 198 | 198 | 1308  |

|        |     |     |     |      |     |     |     |      |
|--------|-----|-----|-----|------|-----|-----|-----|------|
| Ffit_2 | 948 | 210 | 234 | 1392 | 470 | 159 | 160 | 789  |
| Ffit_3 | 155 | 32  | 36  | 223  | 156 | 39  | 37  | 232  |
| Ffit_4 | 245 | 43  | 49  | 337  | 265 | 51  | 49  | 365  |
| Ffit_5 | 389 | 102 | 130 | 621  | 689 | 210 | 267 | 1166 |
| Ffit_6 | 34  | 15  | 12  | 61   | 49  | 19  | 12  | 80   |

This research work developed an AI-based CoV19 feature prediction model, which has been tested with original and data-augmented datasets. All the scan images are resized based on the feature size of 256x256 pixels with a change in aspect ratio, compare to the original. Figure 2 (a) and (b) illustrate the scan images of CoV-19 contaminated patient and a healthy patient, respectively. The CoV19 detection model is a combination of the proposed model FVisL-CoV19 and VGG16, through AI and they are used to detect CoV19 using scan images. Figures 2 (c) and (d) illustrate the scan images of CoV-19 infected patient and a healthy patient, respectively. The collected scans of scans were split into a training and a testing set, which are illustrated in table 2.



## B. Performance Evaluation:

The recommended model is to identify CoV19 and non-CoV19 samples based on the performance metrics illustrated in Table 3: Prediction metrics.

Table 3: Performance metric for prediction

| Performance metric | Definition                        |
|--------------------|-----------------------------------|
| True -tive (TN)    | Appropriately identified as -tive |
| True +tive (TP)    | Appropriately identified as +tive |
| False -tive (FN)   | Imperfectly identified as -tive   |
| False +tive (FP)   | Imperfectly identified as +tive   |

Based on Table 3, the evaluation metrics formulae are illustrated in Table 4.

Table 4: Evaluation metrics

| Evaluation metric    | Definition                          |
|----------------------|-------------------------------------|
| A (in %) (Accuracy)  | $\frac{TP + TN}{TP + FP + TN + FN}$ |
| P (in %) (Precision) | $\frac{TP}{TP + FP}$                |
| R (Recall)           | $\frac{TP}{TP + FN}$                |
| F1 (F1 score)        | $2 * \frac{R * P}{R + P}$           |

Table 5: Assessment of the New Models' Performance Compared with previous one

| Method                | P    | R    | F1   | A    |
|-----------------------|------|------|------|------|
| InceptionV3 [16]      | 0.83 | 0.85 | 0.84 | 0.88 |
| DensNet [17]          | 0.92 | 0.91 | 0.92 | 0.91 |
| Proposed <b>FfitL</b> | 0.94 | 0.93 | 0.94 | 0.93 |

Table 5 suggests that each parameter of the assessment values, specific accuracy, is over 0.88. Therefore, the Proposed FfitL is a methodical model to find COV-19 and non-COV-19 features. In this work, find that the FfitL-CoV19 model performs very well when compared to the others, and we propose it as a replacement for the existing FfitL models.

## V. Conclusions:

In this paper, research is made through the AI techniques for the presence of Covid-19 through the characteristic evaluation that has been proposed. The purpose has been to find out the maximum dependable technique and to enhance the model-primarily based total evaluation. After visual-type classification, the features were determined that the proposed classifier offers higher effects through 94% precision, 93% recall, 94% f1-score, and with 93% accuracy. Future work will consider the usage of the AI multiplicative nature of COV-19 infections with the aid of using building a structural model to recognize the infectious disease-spreading nature of COV-19 and measures to slow down the spread of COV-19.

## VI. References

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