RISK PREDICTION OF CRITICAL VITAL SIGNS FOR ICU PATIENTS USING RECURRENT NEURAL NETWORK

A Thesis

Presented to

The Faculty of the Department of Computer Science
California State University, Los Angeles

In Partial Fulfillment

of the Requirements for the Degree

Master of Science

in

Computer Science

By

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December 2019

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December 2019

ABSTRACT

Risk Prediction of Critical Vital Signs for ICU Patients Using Recurrent Neural Network

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Monitoring vital signs for Intensive Care Unit (ICU) patients is an absolute necessity to help assess the general physical health. In this research, we use machine learning to make a classification forecast that uses continuous ICU vital signs measurements to predict whether the vital signs of the next hour would reach the critical value or not. With the early warning, nurses and doctors can prevent emergency situations that may cause organ dysfunction or even death before it is too late.

In this study, the data includes vital sign measurements, laboratory test results, procedures, medications collected from over 40,000 patients. After data preprocessing, bias data balancing, feature extraction, and feature selection, we have a clean dataset with informative and discriminating features. Then, various machine learning algorithms including Random Forest, XGBoost, Artificial Neural Networks (ANN), and LSTM were developed to predict critical vital signs of ICU patients 1-hour beforehand. We particularly developed predictive models to predict Heart Rate, Blood Oxygen Level (SpO2), Mean Arterial Pressure (MAP), Respiratory Rate (RR), Systolic Blood Pressure (SBP). The results demonstrated the accuracy of the developed methods.

ACKNOWLEDGMENTS

I would like to appreciate MIT Laboratory for Computational Physiology and collaborating research groups for providing valuable data. I would also like to thank Dr. Mohammad Pourhomayoun Ph.D. for guiding me improving the performance of this thesis. And, I want to thank Dr. David Ray Chang M.D. for giving the medical knowledge and suggestions.

TABLE OF CONTENTS

| Abstract | iv |
|---------------------------------|------|
| Acknowledgments | V |
| List of Tables | vii |
| List of Figures | viii |
| Chapter | |
| 1. Introduction | 1 |
| Purpose of Project | 2 |
| 2. The Dataset | 3 |
| Features and Label | 3 |
| Combine Duplicate Measurements | 4 |
| Timeline Alignment | 5 |
| Remove Error Measurements | 6 |
| 3. Types of Dataset | 8 |
| Situations in ICU | 8 |
| Bias of Data | 9 |
| 4. Feature Extraction | 11 |
| 5. Deep Learning Method | 12 |
| Artificial Neural Network (ANN) | 12 |
| Recurrent Neural Network (RNN) | 13 |
| Long Short-Term Memory (LSTM) | 13 |
| 6. Conclusion | 15 |
| References | 17 |

LIST OF TABLES

| Table |
|-------|
|-------|

| 1. | TABLE 1. STAGES OF HYPOVOLEMIC SHOCK | 1 |
|----|---------------------------------------|----|
| 2. | TABLE 2. SOME VITAL SIGN MEASUREMENTS | 4 |
| 3. | TABLE 3. BEST RESULTS FOR VITAL SIGNS | 15 |

LIST OF FIGURES

| Figure 1. | Item Table of Systolic Blood Pressure | 5 |
|------------|---|---|
| Figure 2. | Non-hourly measurement raw data | 5 |
| Figure 3. | Continuously Measured Respiratory Rate of Patient 6 | 6 |
| Figure 4. | Error measurements of SpO ₂ | 7 |
| Figure 5. | 9 Situations in ICU | 8 |
| Figure 6. | Current Normal Situations in ICU | 9 |
| Figure 7. | Balanced Training Dataset | 0 |
| Figure 8. | Concept of Feature Extraction | 1 |
| Figure 9. | Structure of Artificial Neural Network (ANN) | 2 |
| Figure 10. | Structure of Recurrent Neural Network (RNN) | 3 |
| Figure 11. | Features and Target in LSTM model | 4 |
| Figure 12. | AUC Results from all Algorithms | 6 |
| Figure 13. | ROC curves for RNN-LSTM model | 6 |

Introduction

In an intensive care unit (ICU), vital signs indicate the status of the patient's life-sustaining functions. If any of the vital signs has reached the critical value, the patient is in danger and needs immediate help from nurses or doctors. Prolonged hypoperfusion in critically ill patients leads to multiple organ failure, which further increases mortality. Hypoperfusion usually presents as unstable vital signs, which is instantly warned via the ICU alarm [2],[3].

Frequently in the critically ill setting, the alarm from the ICU monitor is too slow for intervention. For example, the ICU monitor will only alarm when tachycardia occurs in stage 2 hypovolemic shock patients with blood loss 15-30%, or when hypotension occurs in stage 3 with blood loss over 30%. The optimal timing of treatment should be stage 1 when the patient only has blood loss under 15%, but the vital signs will mostly be normal at this stage [4].

TABLE 1. STAGES OF HYPOVOLEMIC SHOCK

| | Stage I | Stage II | Stage III | Stage IV |
|----------------|---------|---------------------------|-------------------|------------------|
| Blood Loss | <750 | 750 - 1500 | 1500 - 2000 | > 2000 |
| % Blood Vol. | <15% | 15 – 30% | 30 – 40% | > 40% |
| Heart Rates | <100 | >100 | >120 | >140 |
| Blood Pressure | Normal | Increased diastolic BP | Systolic BP < 100 | Systolic BP < 70 |
| Resp. Rate | 14 - 20 | 20 – 30 | 30 – 40 | > 40 |
| Mental Status | Normal | Anxious | Confused | Lethargic |

Purpose of Project

An early prediction could help doctors and nurses be aware of critical vital signs in advance and prevent the patient from deteriorating. By using an accurate predictive model, in a list of patients with normal vital signs, doctors and nurses will know which patients to look into first. Thus, the medical team will have more time for diagnosis and intervention.

The rapid advances in data science, body sensors, and artificial intelligence have led to the development of effective patient monitoring and data-driven analytics systems that allow for gathering information from patients and analyze it to predict health conditions [8]. These systems have shown potential effectiveness in decreasing healthcare costs and reducing morbidity and mortality [8]-[22].

In this paper, we develop an accurate predictive model using several machine learning algorithms including Recurrent Neural Network (RNN), Long Short-Term Memory (LSTM) networks to predict critical vital signs of ICU patients 1-hour beforehand.

The Dataset

The data of this paper, MIMIC-III (Medical Information Mart for Intensive Care III) [5], includes patient general information, vital sign measurements, laboratory test results, procedures, medications, and mortality collected from over 40,000 patients between 2001 and 2012. In total, we collected 1,323,067 (hours) data samples from all patients after the data preprocessing.

Features and Label

In general ICU, if primary vital signs are out of normal range, then ICU monitors would warn the nurses that patients have abnormal situations. Thus, we want to predict whether the vital signs of the next hour will change to abnormal (above upper bound, below lower bound) or still stays in the normal range, which are our two target classifications. In other word, the goal is to predict the transition from currently normal status to future abnormal status.

Table 2 shows the list of primary vital sign measurements and its normal range based on the National Early Warning Score (NEWS) and AHA ACLS guidelines [6], [7]. We are using 6 vital signs which are the features in 6 different individual models to predict 0 as normal or 1 as abnormal for each vital sign.

TABLE 2. SOME VITAL SIGN MEASUREMENTS [6], [7]

| Feature | Normal Range |
|--|------------------------------|
| Heart Rate (HR) | 50 to 130 (beats per minute) |
| Blood Oxygen Level (SpO ₂) | 90% to 100% |
| Mean Arterial Pressure (MAP) | 65 to 110 (mm Hg) |
| Respiratory Rate (RR) | 8 to 30 (Breaths per minute) |
| Systolic Blood Pressure (SBP) | 90 to 160 (mm Hg) |
| Diastolic Blood Pressure (DBP) | 60 to 110 (mm Hg) |

Combine Duplicate Measurements

Because this 45 Gig Big Dataset is collected from over 600 hospitals and each hospital using different LABEL name, we have the same measurements with different item id. As shown in Fig. 2, we can notice that "abp [systolic]" is an abbreviation of "arterial bp [systolic]", but the ITEMID is different. We also can notice that blood pressure has been measured from different part of body.

In order to deal with this situation, we combine all the vital sign measurements if they represent the same. Thus, "abp [systolic]" and "arterial bp [systolic]" would have the same ITEMID.

| | ROW_ID | ITEMID | LABEL | ABBREVIATION | DBSOURCE | LINKSTO | CATEGORY | UNITNAME | PARAM_TYPE | CONCEPTID |
|------|--------|--------|------------------------------------|--------------|----------|-------------|----------|----------|------------|-----------|
| 295 | 32 | 6 | abp [systolic] | NaN | carevue | chartevents | NaN | NaN | NaN | NaN |
| 320 | 57 | 51 | arterial bp [systolic] | NaN | carevue | chartevents | NaN | NaN | NaN | NaN |
| 671 | 408 | 442 | manual bp [systolic] | NaN | carevue | chartevents | NaN | NaN | NaN | NaN |
| 682 | 419 | 455 | nbp [systolic] | NaN | carevue | chartevents | NaN | NaN | NaN | NaN |
| 705 | 442 | 480 | orthostat bp sitting [systolic] | NaN | carevue | chartevents | NaN | NaN | NaN | NaN |
| 707 | 444 | 482 | orthostatbp standing [systolic] | NaN | carevue | chartevents | NaN | NaN | NaN | NaN |
| 709 | 446 | 484 | orthostatic bp lying [systolic] | NaN | carevue | chartevents | NaN | NaN | NaN | NaN |
| 715 | 452 | 492 | pap [systolic] | NaN | carevue | chartevents | NaN | NaN | NaN | NaN |
| 1437 | 618 | 666 | systolic unloading | NaN | carevue | chartevents | NaN | NaN | NaN | NaN |
| 1748 | 929 | 3313 | bp cuff [systolic] | NaN | carevue | chartevents | NaN | NaN | NaN | NaN |
| 1750 | 931 | 3315 | bp left arm [systolic] | NaN | carevue | chartevents | NaN | NaN | NaN | NaN |
| 1752 | 933 | 3317 | bp left leg [systolic] | NaN | carevue | chartevents | NaN | NaN | NaN | NaN |
| 1754 | 935 | 3319 | bp pal [systolic] | NaN | carevue | chartevents | NaN | NaN | NaN | NaN |
| 1756 | 937 | 3321 | bp right arm [systolic] | NaN | carevue | chartevents | NaN | NaN | NaN | NaN |
| 1758 | 939 | 3323 | bp right leg [systolic] | NaN | carevue | chartevents | NaN | NaN | NaN | NaN |
| 1760 | 941 | 3325 | bp uac [systolic] | NaN | carevue | chartevents | NaN | NaN | NaN | NaN |
| | | | | | | | | | | |

Fig. 1. Item Table of Systolic Blood Pressure

Timeline Alignment

Although the raw data is collected by ICU monitors, the timestamps of all measurements are not unified. Since more than 80% of the raw data are measured on hourly basis, we fixed the data collection rate to one sample per hour.

| : | | SUBJECT_ID | HADM_ID | ICUSTAY_ID | ITEMID | CHARTTIME | STORETIME | CGID | VALUE | VALUENUM | VALUEUOM | WARNING |
|---|----------|------------|---------|------------|--------|------------------------|------------------------|---------|-------|----------|----------|---------|
| | 34061542 | 3 | 145834 | 211552.0 | 807 | 2101-10-21 06:00:00 | 2101-10-21 05:46:00 | 21570.0 | 306 | 306.0 | NaN | NaN |
| | 34028522 | 4 | 185777 | 294638.0 | 807 | 2191-03-16 08:00:00 | 2191-03-16 07:49:00 | 17144.0 | 266 | 266.0 | NaN | NaN |
| | 34297591 | 6 | 107064 | 228232.0 | 807 | 2175-05-30 21:00:00 | 2175-05-30 21:54:00 | 15443.0 | 254 | 254.0 | NaN | NaN |
| | 34086330 | 9 | 150750 | 220597.0 | 807 | 2149-11-13 12:00:00 | 2149-11-13 19:10:00 | 20855.0 | 212 | 212.0 | NaN | NaN |
| | 34062720 | 13 | 143045 | 263738.0 | 807 | 2167-01-08 23:30:00 | 2167-01-08 23:21:00 | 21205.0 | 205 | 205.0 | NaN | NaN |
| | 34049478 | 18 | 188822 | 298129.0 | 807 | 2167-10-03 11:00:00 | 2167-10-03 11:01:00 | 19667.0 | 243 | 243.0 | NaN | NaN |
| | 34035552 | 21 | 111970 | 216859.0 | 807 | 2135-02-04 06:00:00 | 2135-02-04 06:05:00 | 15907.0 | 200 | 200.0 | NaN | NaN |
| | 34033889 | 25 | 129635 | 203487.0 | 807 | 2160-11-02 04:45:00 | 2160-11-02 05:23:00 | 18654.0 | 353 | 353.0 | NaN | NaN |
| | 34406341 | 43 | 146828 | 225852.0 | 807 | 2186-10-04 07:00:00 | 2186-10-04 07:29:00 | 17435.0 | 210 | 210.0 | NaN | NaN |
| | 34510579 | 56 | 181711 | 275642.0 | 807 | 2104-01-03 16:00:00 | 2104-01-03 16:19:00 | 21347.0 | 201 | 201.0 | NaN | NaN |
| | 34486585 | 62 | 116009 | 216609.0 | 807 | 2113-02-15 22:00:00 | 2113-02-15 22:55:00 | 15023.0 | 221 | 221.0 | NaN | NaN |
| | 34483962 | 68 | 108329 | 272667.0 | 807 | 2174-01-08 14:00:00 | 2174-01-08 14:05:00 | 14375.0 | 212 | 212.0 | NaN | NaN |
| | | | | | | | | | | | | |

Fig. 2. Non-hourly measurement raw data

Remove Error Measurements

After visualizing the features, we found that some of the measurements of ICU monitor was incorrect. As shown in Fig. 3, for patient 6, respiratory rate has suddenly fallen to 0 between 0 to 10 hours. This was an abnormal situation in ICU, so we considered it as an error from ICU monitor. Thus, we removed all the sudden 0 value of RR. As shown in Fig. 4, the percentage of oxygen in blood (SpO₂) would not be over 100%. We think this was a kind of typo from nurses. However, after talking to the ICU nurses, they said if the measurements values are error or incorrect in ICU, then they would record above 100%. Thus, we remove over 100% value of SpO₂ as well.

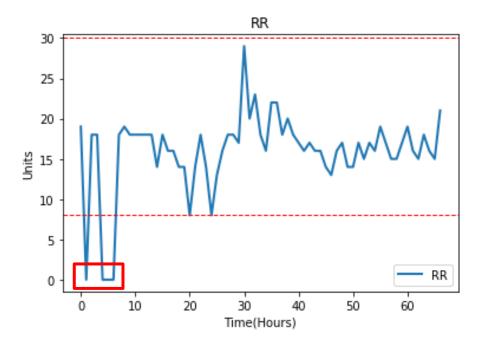


Fig. 3. Continuously Measured Respiratory Rate of Patient 6

| | SUBJECT_ID | ICUSTAY_ID | CHARTTIME | heart rate | spo2 | arterial bp mean | respiratory rate | arterial bp [diastolic] | arterial bp [systolic] |
|---------|------------|------------|------------------------|---------------|--------|------------------------|---------------------|-------------------------------|------------------------------|
| 112892 | 28187 | 297092.0 | 2109-04-10 13:00:00 | 101.0 | 101.00 | 76.0 | 31.0 | 60.0 | 107.0 |
| 698209 | 13183 | 297445.0 | 2117-06-11 13:00:00 | 56.0 | 101.00 | 83.0 | 28.0 | 63.0 | 131.0 |
| 874907 | 17699 | 211512.0 | 2106-10-26 19:00:00 | 96.0 | 100.25 | 84.0 | 10.0 | 61.0 | 140.0 |
| 1068157 | 22596 | 270858.0 | 2133-06-02 20:00:00 | 83.0 | 102.00 | 104.0 | 77.0 | 77.0 | 124.0 |

Fig. 4. Error measurements of SpO_2

Types of Dataset

Situations in ICU

In ICU, we will have 9 different situations. As Shown in fig. 5, current value of vital sign could be normal, low (below lower bound of normal range), and high (above upper bound of normal range), and the next hour value could be normal, low, high as well. Thus, we would get 9 different situations in ICU. However, we only want to predict for the normal value of current hour of vital sign because the main purpose of this study is to warn the nurses and the doctors that this patient is going to deteriorate in this hour. As shown in fig 6, we only need to consider about 3 situations in ICU.

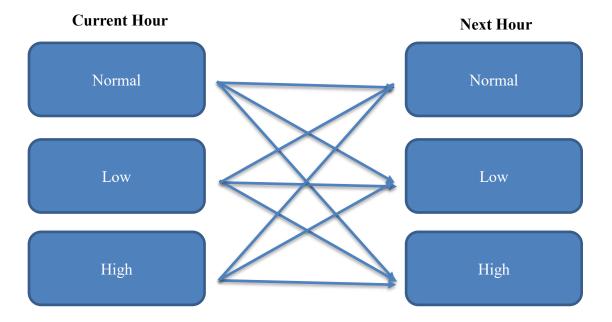


Fig. 5. 9 Situations in ICU

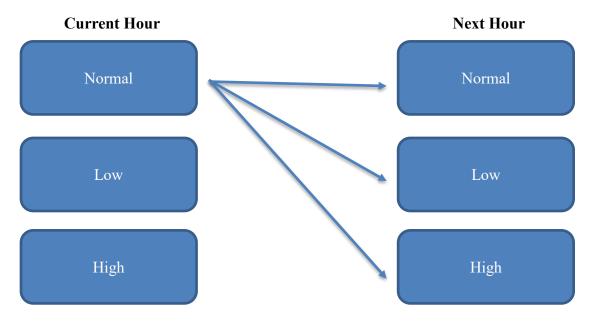


Fig. 6. Current Normal Situations in ICU

Bias of Data

Bias is a common machine learning problem, especially in the medical domain. Machine learning would keep predicting the biased result because the occurrence rate of the healthy state is much higher than the diseased. In our dataset, 99% of HR vital sign is in the normal range and the next hour is still in the normal range as well. The machine learning algorithm trained by the original unbalanced dataset would only predict "normal" for the next hour and would still have a 99% accuracy. In order to avoid this bias problem, we first filtered out all currently abnormal data samples because the goal is to predict the transition from currently normal status to future abnormal status (please note that the patients whose vital signs are currently in abnormal status are not of our prediction interest. We would like to predict whether a patient who is currently in normal situation will change to abnormal range in the next hour or still stays in the normal range).

Secondly, we balanced the binary-target classification in the Training dataset. Thus, we have 50% next hour data within the normal range, 50% next hour data greater than the upper bound and less than the lower bound in the training dataset. (Fig. 7).

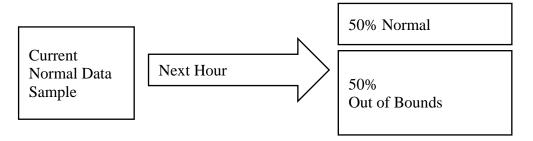


Fig. 7. Balanced Training Dataset

Feature Extraction

As shown in fig.8, with the hourly-measured dataset, we extracted new features from the past 3- and 5-hours window to increase the accuracy, such as the mean and standard deviation of every measurement of the past 3 hours and the past 5 hours. These features can perform how vital signs varied in over the past few hours which are relevant to the next hour value. We derive 24 features from the 3-hour window and 5-hour window.

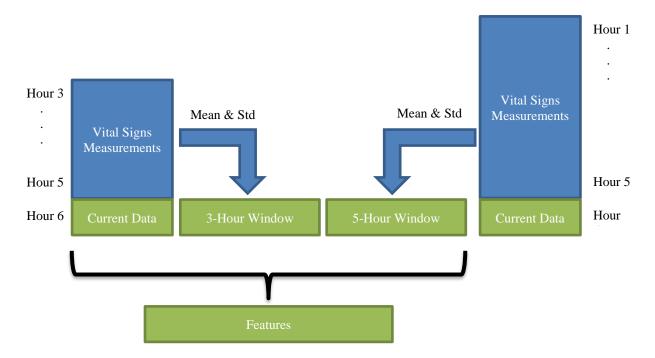


Fig. 8. Concept of Feature Extraction

Deep Learning Method

In this study, we have used several classification machine learning algorithms, such as Random Forest, XGBoost, Artificial Neural Net (ANN), and Recurrent Neural Network (RNN), LSTM. However, we have achieved the best accuracy results with deep learning method, RNN-LSTM.

Artificial Neural Network (ANN)

Artificial Neural Network (ANN) is a computing system like biological neural networks. As shown in fig. 9, the input X are the features, the second layer is neurons, and the output Y as the prediction [26]. The connections between layers are the weights of each nodes. Unlike traditional machine learning, with the calculations of the weights by the neurons, prediction would be more accurate than machine learning.

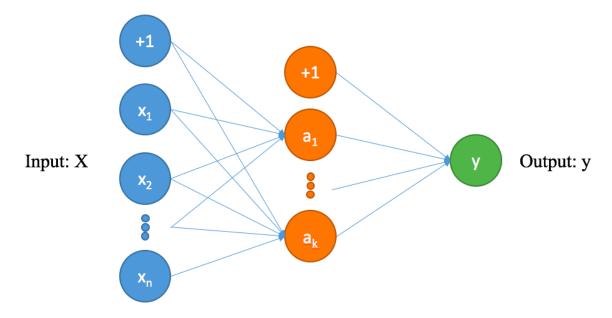


Fig. 9. Structure of Artificial Neural Network (ANN) [26]

Recurrent Neural Network (RNN)

Humans don't start over their thinking when they are making decision. They would make the decision based on their knowledge or experience. For example, you want to predict who would be killed at the next episode of Game of Thrones. It is difficult for traditional ANN because ANN is unclear the time series of all the events.

One of the most effective algorithms for this application is deep RNN due to the strong temporal correlation in the data. As shown in fig. 10, RNN has shown strong potential on forecasting outcomes of time-series or sequential data [24]. Unlike traditional feedforward neural networks, RNN includes feedback connections. It allows the algorithm to memorize historical data and consequently, use a sequence of data samples to make more accurate predictions when there is a temporal correlation in the input data [23].

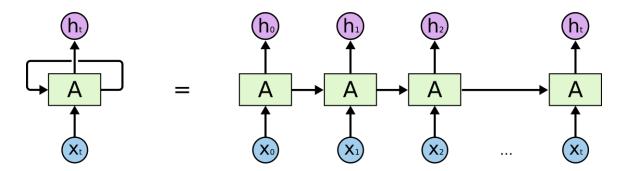


Fig. 10. Structure of Recurrent Neural Network (RNN) [24]

Long Short-Term Memory (LSTM)

A serious problem in training every deep neural network including RNN is vanishing gradient. Vanishing gradient happens when the gradient and, consequently, the

network weight correction in the training process (e.g. backpropagation), tends to be very close to zero, especially when we move backward towards the front layers. It makes the training process very slow. This problem is even more serious for RNN with many timesteps.

The Long Short-Term Memory (LSTM) structure can help solve the vanishing gradient problem in traditional RNN [25]. An LSTM unit usually includes a cell, an input gate, an output gate and a forget gate. In this project, we will design and train RNNs with LSTM units. In this approach, the temporal correlation will be taken into account by the memory of the LSTM and the sequential nature of the input data.

As shown in fig. 11, the input would be current hour and the past 5 hours with extractions of the features, and the output would be our target hour 7.

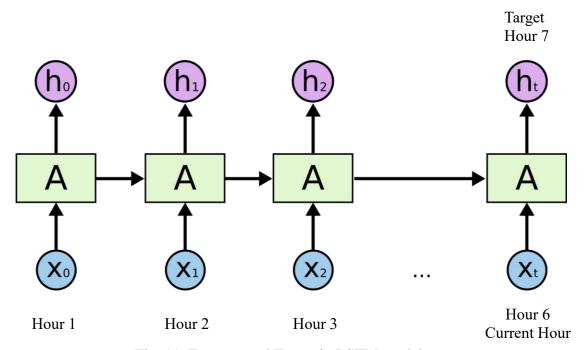


Fig. 11. Features and Target in LSTM model

Conclusion

In this paper, we develop accurate predictive models using various machine learning algorithms including deep RNN with LSTM units to predict critical vital signs of ICU patients 1-hour beforehand. This system can significantly help doctors and nurses be aware of critical vital signs in advance and prevent the patient from deteriorating.

The predictive models were trained and tested on a big dataset including the vital sign measurements, laboratory test results, procedures, medications collected from over 40,000 patients. We randomly split the data samples into training (70%) and testing (30%) datasets.

As shown in Chart I and Table II, for all the vital signs, the RNN-LSTM has the best score of Area Under the Curve (AUC) based on the True Positive Rate (TPR) and the False Positive Rate (FPR) plane. Fig. 2 demonstrates ROC curves for RNN-LSTM model.

TABLE 3. BEST RESULTS FOR VITAL SIGNS

| Vital signs | AUC Score |
|--|-----------|
| Heart Rate (HR) | 91.3% |
| Blood Oxygen Level (SpO ₂) | 80.1% |
| Mean Arterial Pressure (MAP) | 75.0% |
| Respiratory Rate (RR) | 82.2% |
| Systolic Blood Pressure (SBP) | 78.7% |

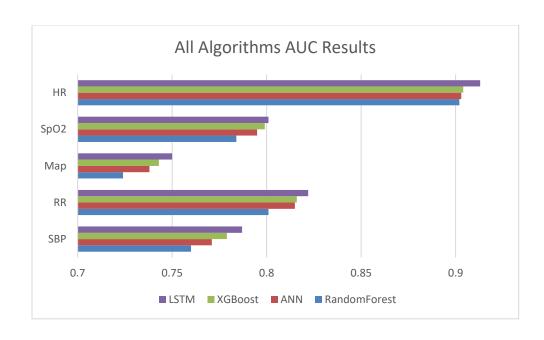


Fig. 12. AUC Results from all Algorithms

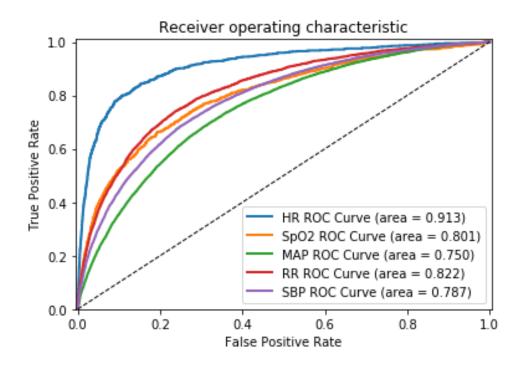


Fig. 13. ROC curves for RNN-LSTM model

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