Forecasting Thresholds Alarms in Medical Patient Monitors using Time Series Models

Jonas Chromik¹[®]^a, Bjarne Pfitzner¹[®]^b, Nina Ihde¹[®]^c, Marius Michaelis¹[®]^d, Denise Schmidt¹[®]^e, Sophie Anne Ines Klopfenstein²[®]^f, Akira-Sebastian Poncette²[®]^g, Felix Balzer²[®]^h

and Bert Arnrich¹^[D]

¹Hasso Plattner Institute, University of Potsdam, Germany ²Charité – Universitätsmedizin Berlin, Berlin, Germany

Keywords: Patient Monitor Alarm, Medical Alarm, Intensive Care Unit, Vital Parameter, Time Series Forecasting, Alarm Forecasting, Alarm Fatigue.

Abstract: Too many alarms are a persistent problem in today's intensive care medicine leading to alarm desensitisation and alarm fatigue. This puts patients and staff at risk. We propose a forecasting strategy for threshold alarms in patient monitors in order to replace alarms that are actionable right now with scheduled tasks in an attempt to remove the urgency from the situation. Therefore, we employ both statistical and machine learning models for time series forecasting and apply these models to vital parameter data such as blood pressure, heart rate, and oxygen saturation. The results are promising, although impaired by low and non-constant sampling frequencies of the time series data in use. The combination of a GRU model with medium-resampled data shows the best performance for most types of alarms. However, higher time resolution and constant sampling frequencies are needed in order to meaningfully evaluate our approach.

SCIENCE AND TECHNOLOGY PUBLICATIONS

1 INTRODUCTION

Alarm fatigue is a persisting problem in today's intensive care medicine with negative outcomes for patients and staff (Cvach, 2012). Although the problem is well understood from a medical point of view, there is no sufficient technical solution to the alarm fatigue yet. Among the alarms produced by medical patient monitors, threshold alarms are of particular interest. Alarms, in general, are supposed to express acute critical events that need immediate attention. This is, for example, the case with arrhythmia alarms. Threshold alarms, however, do not necessarily result from

- ^a https://orcid.org/0000-0002-5709-4381
- ^b https://orcid.org/0000-0001-7824-8872
- ^c https://orcid.org/0000-0001-5776-3322
- ^d https://orcid.org/0000-0002-6437-7152
- e https://orcid.org/0000-0002-6299-0738
- ^f https://orcid.org/0000-0002-8470-2258
- g https://orcid.org/0000-0003-4627-7016
- h https://orcid.org/0000-0003-1575-2056
- ⁱ https://orcid.org/0000-0001-8380-7667

an acute event but can be the result of a continued trend as we learned from a contextual inquiry at an intensive care unit (ICU).

In this paper, we want to forecast the foreseeable share of threshold alarms in order to transform these alarms into scheduled tasks, thus removing the urgency of the situation. Rather than having an alarm that has to be taken care of immediately, we want to present scheduled tasks to the medical staff. For example: "In approximately one hour the blood pressure of the patient will rise above the high threshold. During the next hour, some member of staff should take care of this issue." This is especially relevant since the majority of audible and actionable alarms are threshold alarms (Drew et al., 2014).

Forecasting threshold alarms is done in this work by means of time series models, such as autoregressive integrated moving average (ARIMA) models and recurrent neural networks (RNNs). As data source we chose the MIMIC-III database (Johnson et al., 2016) since this is – to the best of our knowledge – the only clinical database containing data on patient monitor alarms. Our approach is optimised for high speci-

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Chromik, J., Pfitzner, B., Ihde, N., Michaelis, M., Schmidt, D., Klopfenstein, S., Poncette, A., Balzer, F. and Arnrich, B. Forecasting Thresholds Alarms in Medical Patient Monitors using Time Series Models.

DOI: 10.5220/0010767300003123

In Proceedings of the 15th International Joint Conference on Biomedical Engineering Systems and Technologies (BIOSTEC 2022) - Volume 5: HEALTHINF, pages 26-34 ISBN: 978-989-758-552-4; ISSN: 2184-4305

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ficity rather than high sensitivity since we acknowledge that not all threshold alarms are the continuation of a prolonged trend and hence foreseeable. We want to avoid exacerbating the problem of alarm fatigue by false positives of our approach.

The rest of this work is structured as follows: In section 2 we describe the data and models we use in this work. In section 3 we present the results we achieve with our approach. In section 4 we discuss these results. Finally, in section 5 we conclude our work.

2 MATERIALS & METHODS

In order to forecast threshold alarms in medical patient monitors, we need both data on said threshold alarms and means to forecast the corresponding vital parameter. As data source, we use the MIMIC-III clinical database as we describe in section 2.1. As forecasting methods, we compare a variety of time series models as we describe in section 2.2.

2.1 Materials

The MIMIC-III database contains 26 tables providing a wide range of information on the events at the ICUs of Beth Israel Deaconess Medical Center. For our use case, however, only the CHARTEVENTS table is of interest. This table contains, among others, measured values and alarm thresholds of a variety of vital parameters such as heart rate (HR), respiratory minute volume (MV), non-invasively measured systolic blood pressure (NBPs), respiratory rate (RR), and peripheral blood oxygen saturation (SpO₂). For forecasting, we require time series with sufficiently high and relatively stable sampling frequencies over an extended period of time. This is not the case for all vital parameters. Thus, we chose to use only HR, NBP_s, and SpO₂ as these vital parameters satisfy the aforementioned requirements. Specifically, we are interested in the data items listed in table 1.

2.2 Methods

We want to apply time series forecasting on the vital parameter data described in section 2.1 in order to achieve our ultimate goal of forecasting threshold alarm events. To do this, we require methods for forecasting time series. For this work, we decided to employ two fundamental approaches: statistical models and machine learning models.

Statistical models aim at forecasting the future of a time series with only the time series itself as prior

Table 1: Complete list of ITEMIDs in the CHARTEVENTS table that are relevant for this work.

ITEMID	Label
220045	HR
220046	HR Alarm - High
220047	HR Alarm - Low
220179	NBP _s
223751	NBP _s Alarm - High
223752	NBP _s Alarm - Low
220277	SpO ₂
223769	SpO ₂ Alarm - High
223770	SpO ₂ Alarm - Low

knowledge. Specifically, we are using the ARIMA model and its variation additionally featuring exogenous variables (ARIMAX).

Machine learning (ML) models learn from training data and can then be applied to previously unseen test data. In our case, we employ RNNs trained with 80% of the available vital parameter time series to forecast on the remaining 20% of vital parameter time series. Specifically, we are using vanilla RNNs, gated recurrent units (GRUs), and long short-term memory neural networks (LSTMs) because variations of recurrent neural networks is what is usually used on timeseries data, e.g. in (Mussumeci and Coelho, 2020), (Pathan et al., 2020), and (Dai et al., 2021).

We frame the problem as a regression problem, i.e. forecasting the vital parameter, instead of a classification problem, i.e. predicting whether an alarm will occur or not, because we want to keep the ML approach as close to the ARIMA approach as possible to ensure comparability. For the same reason, we also do not provide additional information such as age or sex to the ML models.

Resampling and Chunking. Both statistical and machine learning models have in common that they require constant sampling frequencies which are not always given in medical databases. We address this issue by resampling the vital parameter measurement from the MIMIC-III database to one sample per hour $(f_s = 1 \text{ h}^{-1})$ which is close to the database's original sampling frequency present in most cases.

For resampling, we employ three different strategies in order to fuse samples together: Minimum resampling, maximum resampling, and median resampling. Thus, we create three distinct but related time series which can be used for forecasting.

Furthermore, we use a chunking strategy. We noticed that there are gaps in the time series, i.e. extended periods of time where there are no data points. We assume, that this is due to the patient being in a different ward, surgery, or some other procedure. These missing data pose a difficulty for resampling. Hence, we subdivide the data of a patient's ICU stay along the data gaps into multiple chunks and operate only on these chunks throughout the rest of our work.

By subdividing the patients' data into chunks and treating these chunks as distinct time series, we circumvent a missing data issue. In practice, this implies that whenever a period of missing data arises, the model has to re-learn and cannot build upon the data from the prior chunk. Nevertheless, we reason that this approach makes sense because we cannot know what happened in the period of missing data. For example, when the period of missing data was caused by a surgical procedure, the patient might be in a completely different condition after the surgery than before. Consequently having a completely different vital parameter distribution that can not be associated with the period before the surgery.

Experiment Setup. Regardless of whether the model in use is a statistical model or a ML model, we always employ the same experiment setup: We use 12 or 30 timesteps (lags) equivalent to 12 or 30 hours of vital parameter data as input for the model. We chose these specific periods because we wanted to compare performances for a rather short and a rather long observation. The 12 hours period primarily aims at providing clinicians with alarm forecasts in a timely manner. With this approach, forecasts are provided after the patient spent half a day at the ICU. In contrast, the 30 hours time frame aims at sufficiently spanning a complete cycle of the circadian rhythm hence giving a more holistic picture of the patient's vital parameter distribution.

The model produces a forecasted value for the hour following the input lags. We compare the forecasted value to the currently active high or low alarm threshold. If the value is above the high threshold or below the low threshold, a respective threshold alarm is forecasted. Otherwise, no alarm is forecasted. Subsequently, we compare the forecast with the actual situation, i.e. whether there was actually a threshold alarm triggered by the actual vital parameter measurement.

This common approach is used for all models we evaluated. However, details differ since there are conceptual differences between the models. The most striking one is that ML models require dedicated training data while statistical models learn only on the given input sequence. Hence, we describe the concrete experiment setups for specific models in the following. **Statistical Models.** We use both ARIMA and ARI-MAX models as can be seen in Table 2. The ARIMA models are used with the median and minimum or maximum resampled time series. For ARIMAX we use the maximum resampled time series for forecasting high threshold alarms and the minimum resampled time series for forecasting the low threshold alarms. For both ARIMAX cases, the median resampled time series is used as exogenous series. All three described model setups are processed with a train size of 12 lags or 30 lags, respectively, which determines the minimum length of chunks required.

Table 2: Complete list of IDs for ARIMA and ARIMAX models.

Train Size	Model Type	Endog.
12	ARIMA	Median
12	ARIMA	Min/Max
12	ARIMAX	Min/Max
30	ARIMA	Median
30	ARIMA	Min/Max
30	ARIMAX	Min/Max
	Train Size 12 12 12 30 30 30	Train SizeModel Type12ARIMA12ARIMA12ARIMAX30ARIMA30ARIMA30ARIMA

Machine Learning Models. Here, we also combine different models with different resampling strategies. As model types we chose vanilla RNNs, GRUs, and LSTMs. As with statistical models, we have one setup that is equivalent to the ARIMA setup where we train and forecast only with the median resampled time series. In another setup which is equivalent to the ARIMAX approach, we use the maximum resampled time series for forecasting high threshold alarms and the minimum resampled time series for forecasting the low threshold alarms and in both cases the median resampled time series as exogenous series.

To make a prediction for each chunk, we introduce a windowing technique that uses 80% of the data of the respective vital parameter for training and 20% for predicting per window. Thus, a total of five different windows are considered, each predicting different 20% of the chunks ensuring that no previously seen data is used. The entire chunk length is employed for training. The prediction on the pre-trained model starts after 12 timesteps, as already described above. We have chosen this "waiting period" in order to better compare the results with those of the ARIMA(X) approach and because a prediction after 12 data points is in practice more crucial than after 30 data points.

As shown in Table 3, each model type is run not only with and without exogenous input, but also with non-scaled (suffix "n"), standard scaled (suffix "s1") and min-max scaled (suffix "s2") time series.

Equation 1 shows how standard scaling is applied on a series value x. It removes the mean and scales

Table 3: Complete list of IDs for ML models. Standard scaling is indicated by suffix "s1" and min-max scaling by suffix "s2". If no scaling is performed, suffix is "n" for "non-scaled".

Model ID	Scaling	Model Type	Endog.
LS_01_s1	Standard	LSTM	Median
LS_02_s1	Standard	LSTM	Min/Max
GR_01_s1	Standard	GRU	Median
GR_02_s1	Standard	GRU	Min/Max
RN_01_s1	Standard	RNN	Median
RN_02_s1	Standard	RNN	Min/Max
LS_01_s2	Min-Max	LSTM	Median
LS_02_s2	Min-Max	LSTM	Min/Max
GR_01_s2	Min-Max	GRU	Median
GR_02_s2	Min-Max	GRU	Min/Max
RN_01_s2	Min-Max	RNN	Median
RN_02_s2	Min-Max	RNN	Min/Max
LS_01_n	None	LSTM	Median
LS_02_n	None	LSTM	Min/Max
GR_01_n	None	GRU	Median
GR_02_n	None	GRU	Min/Max
RN_01_n	None	RNN	Median
RN_02_n	None	RNN	Min/Max
	•		

$$x_{scaled} = \frac{x - \mu}{\sigma} \tag{1}$$

1: Transformation of series value *x* to scaled series value x_{scaled} using **standard scaling (referred to as s1)**. The same mean μ and standard deviation σ is used for all train series and all prediction series.

the values to unit variance. We performed this scaling method globally, meaning all train and all prediction time series are transformed with the same mean μ and same standard deviation σ . This ensures that equal x values are transformed to equal x_{scaled} values across all available time series.

$$x_{scaled} = \frac{x - min}{max - min} \tag{2}$$

2: Transformation of series value x to scaled series value x_{scaled} using **min-max scaling (referred to as s2)**. *min* and *max* are the minimum and maximum of the available data values. This transformation is performed individually for each series resulting in a 0-1 value range.

In contrast to the first scaling method, min-max scaling transforms each value to a given min-max range. In addition, we performed this transformation with each time series individually and not globally. Equation 2 shows how we applied it to our data resulting in a 0-1 value range.

Evaluation. To evaluate the performance of the different models, we employ an adapted version of the

evaluation formula used in (Clifford et al., 2015). There, the goal was to evaluate models for identifying false cardiac arrhythmia alarms. We adapted the formula to ignore true negatives and flipped the roles of false positives and false negatives to account for the inverse scenario we are facing in this paper. The evaluation formula is shown in eq. (3).

Evaluation Score =
$$\frac{TP}{TP + FN + 5 \cdot FP}$$
 (3)

3: Evaluation score formula, adapted from (Clifford et al., 2015).

True positives, present in the numerator and the denominator, represent alarms that are correctly forecast, i.e. the forecasting model predicts an alarm and the alarm is in fact present.

False positives denote situations where the model predicts an alarm but there is in fact none. False positives are penalised with a factor of five because we want to avoid increasing the alarm load since this would be opposed to our goal of alleviating alarm fatigue.

False negatives, on the other hand, are situations where the model predicts no alarm to occur but there is in fact an alarm in the respective period of time. False negatives are less problematic since we acknowledge that changes in vital parameter measurements can happen abruptly and unforeseeably due to external factors that are not recorded in the data set. False negative, in this case, does not mean that the alarm itself is suppressed but rather that the model did not forecast the alarm. Consequently, the alarm stays an indicator for an acute event that needs urgent action rather than being transformed to a scheduled task.

3 RESULTS

In this section, we show and compare the performances of the employed models in terms of our evaluation metric. We first show the performances of the statistical models. Then, we show the performances of the ML models. Finally, we compare the best performing models from both categories amongst each other.

Statistical Models. Figure 1 shows a performance comparison for all statistical models and all parameters with respect to the train size. We compare training with 12 lags against training with 30 lags. We employ a constant sampling frequency of $f_s = 1 \text{ h}^{-1}$ because MIMIC-III does not allow for higher f_s . Hence,

12 lags are equivalent to 12 hours of ICU stay and 30 lags are equivalent to 30 hours of ICU stay, respectively. Since there are fewer ICU stays lasting up to 30 hours than ICU stays lasting up to 12 hours, there are consequently fewer ICU stays to be considered for the 30 lags train size approach.

Figure 2 compares the statistical models' performance for high alarms against the performance for low alarms and across the different vital parameters. There are striking performance differences amongst parameters and alarm types which will be further discussed in section 4.



Figure 1: Comparison of train sizes for statistical models (ARIMA and ARIMAX). For all parameters and model we compare a train size of 12 lags with a train size of 30 lags both for high alarms (suffix _H) and low alarms (suffix _L).



Figure 2: Comparison of alarm types (high alarm and low alarm) for statistical models across all vital parameters.

ML Models. Figure 3 shows a comparison of the different ML model types, i.e. vanilla RNN, LSTM, and GRU. The plots show no clear superiority of one model type. However, the plots suggest that input variables, alarm type, and especially the scaling methods distinctly influence the models' performance. Hence, we further explore the influence of the scaling method in fig. 4. In this figure, two facts are to be seen. Firstly, scaling seems to have a distinctly negative influence on the model's performance. This is most obvious for the SpO₂ vital parameter but the effect is also present for HR and NBP_s. Secondly, the type of alarm (high alarm or low alarm) appears to also have an influence on the ML models' performances but differently than in the statistical models'

case. For HR, low alarms seem to be forecast slightly more successfully. For NBP_s, high alarms seem to be forecast distinctly more successfully. For SpO₂ the influence of the scaling method is too high to clearly see an effect here.

Figure 5 shows an overall comparison of all ML models and highlights the best performing model for each combination of vital parameter and alarm type. For HR and NBP_s there is one clear best model configuration for each alarm type. For SpO₂, however, there are multiple best models for the high alarm type. In general, the GRU model with median resampling shows the overall best performance according to this figure.



Figure 3: Comparison of ML model types vanilla RNN, LSTM and GRU with different configurations across vital parameters and alarm types (suffix _H for high alarms and suffix _L for low alarms).



Figure 4: Comparison of ML models executed with different scaling methods applied (Standard, Min-Max) or without scaling (None) across vital parameters and alarm types (suffix _H for high alarms and suffix _L for low alarms).

Comparison. Figure 6 compares the best performing statistical and ML models with each other having vital parameter and alarm type as independent variable. Except for low blood pressure alarms, the ML models always outperform the statistical models, most strikingly in the SpO₂ case.

To give a more in-depth view into the model performances, fig. 7 visualises the confusion matrix for the best performing models. There, we can see that the statistical models most prominently produce more false positives than the ML models. Since our evaluation metric penalises false positives more heavily than false negatives, this explains the considerably lower performance of the statistical models in fig. 6.



Figure 5: Selection of best ML models. Models with model type GRU and median resampled chunks as endogenous input variable always perform best (except for high alarm forecasting of SpO₂).

4 DISCUSSION

In this paper, we aimed at forecasting threshold alarms in patient monitors. Therefore, we used the vital parameter and alarm data as provided by the MIMIC-III database and employed a wide variety of forecasting models ranging from statistical models to machine learning methods. In general, the overall forecasting performance is not fully satisfactory regardless of the model in use. However, the results give important insights that can guide further research into this area which we discuss in the following.



0.7

0.6

0.6 0.5 0.4 0.3 0.2

0.1

0.0

HR_Low

BP_High S_pO₂ ARIMA X ML 02_High Figure 6: Comparison of best performing statistical models to best performing ML models. Train Sizes. In fig. 1 we compare different ARIMA

HR

NBPs

HR_High

ARIMA_X

ARIMA_X

ML

ARI ML

and ARIMAX models each of which being evaluated with a train of 12 lags and with a train size of 30 lags respectively. As a general finding, longer train sizes tend to yield better forecasting performance. However, a longer train size also entails that the patient needs to stay in the ICU for a longer period of time before the alarm forecasting model can be used, in our case 12 hours vs. 30 hours. Another approach is to raise the time-resolution of the vital parameter data. With more data points per period of time a larger train size can be achieved in less time. This manifests future research work and investigations into ICU databases featuring a higher time resolution, see section 4.2.

High and Low Alarms. Figure 2 and fig. 5 show that model performance vastly differs between high



Figure 7: Comparison of confusion matrix values (false positives, false negatives, and true positives; not showing true negatives) of best performing statistical models to best performing ML models.

and low alarms even concerning the same vital parameter. Especially for HR and NBP_s, high alarms are generally forecast with higher performance regarding the evaluation score we utilise. Furthermore, peak forecasting performance for high and low threshold alarms is not necessarily achieved by the same model. In the cases of HR and SpO₂ forecasting, the peak evaluation score for high alarms is achieved by a different model than the one achieving peak evaluation performance for the low alarm. Consequently, high and low alarms of the same vital parameter have to be considered nonetheless as distinct forecasting tasks. There is no universal one-fits-all model for threshold alarm forecasting.

Effects of Scaling. Scaling does not seem to have a positive effect on the models' forecasting performance, as is to be seen in fig. 4 and fig. 5. In fact, models using unscaled data consistently exhibit superior or hardly worse performance than their counterparts that do apply scaling methods. An example where min-max scaling works better than no scaling is the prediction of alarms of type low in SpO₂. We have no theory on why this is the case. Hence, this manifests a need for further research, ideally in a simplified forecasting setting (e.g. only forecasting the vital parameter measurement and no alarms yet) and time series having a higher and more consistent time resolution, as provided by HiRID and eICU CRD. This is also described as future work in section 4.2.

Best Performing Models. Figure 6 and fig. 7 show that amongst the best performing models (both statistical and ML), the ML models exhibit a superior performance. This is rooted in the evaluation metric we utilise that penalises false positives heavier than false negatives. In fig. 7 it is to be seen that the statistical models tend to produce a higher quantity of false positives which negatively influences their scoring in fig. 6, especially for HR and SpO₂.

4.1 Limitations

In our efforts to forecast threshold alarms in patient monitors we faced a couple of limitations which we already mentioned previously and which we want to summarise here.

Firstly, the relatively low and partially unstable sampling frequency of the vital parameter measurements poses a problem to our approach. This is twofold: On the one hand, the low amount of samples per period of time forces us to use smaller train sizes since larger train sizes would correspond to ridiculously long ICU stays. Hence, we have only 12 to 30 lags of training for the statistical models. We assume that this reduces the forecasting performance of the models since we were able to show that larger train sizes correspond to better model performance. On the other hand, the large temporal distance (\approx 1h) between the samples prevents the forecasting models from picking up on changes that happen on a more fine-grained (higher) temporal resolution, potentially causing false negatives. Higher temporal resolutions in vital parameter measurements and more detailed and accurate alarm event data are needed in order to successfully build accurate alarm forecasting models. Secondly, our work suggests that each alarm type requires a dedicatedly tuned model and that there is no one-fits-all model for forecasting all types of alarms. Hence, a narrower research focus might be required for example limiting the forecasting task on one type of alarms using a more detailed or even specialised data set.

4.2 Future Work

Forecasting threshold alarms in patient monitors is basically an extension of forecasting vital parameter measurements by not only forecasting the value itself but also comparing the value to the alarm thresholds. We chose the MIMIC-III database because a unique feature of this database is that it contains alarm thresholds. However, if we accept that forecasting vital parameter measurements without accounting for alarms is a valid preliminary goal, other clinical databases are eligible as well. For example, HiRID (Hyland et al., 2020) provides vital parameters measurements with a vastly higher time resolution and eICU CRD (Pollard et al., 2018) even provides such data with a steady sampling frequency of $f_s = \frac{1}{5 \min}$ (one value every five minutes). This is in sharp contrast to MIMIC-III which has varying sampling frequencies leaning towards one value per hour. Using HiRID and eICU CRD might improve the forecasting accuracy for vital parameters and also spare us the resampling step which introduces an additional source of inaccuracies and errors. Such a simplified forecasting setting can then also be used to further investigate the effects of scaling on the forecasting performance.

5 CONCLUSIONS

The contribution of this paper is a first attempt to forecasting threshold alarms in ICU patient monitors. Due to the lack of alarm data having a sufficiently high and consistent sampling frequency, the resulting models are still worthy of improvement and are not yet ready to be applied in clinical practice. However, our results show that the general approach of forecasting threshold alarms through vital parameters principally works and that the model setup used in this work is promising.

ACKNOWLEDGEMENTS

This work was partially carried out within the INALO project. INALO is a cooperation project between AICURA medical GmbH, Charité – Universitätsmedizin Berlin, idalab GmbH, and Hasso Plattner Institute. INALO is funded by the German Federal Ministry of Education and Research under grant 16SV8559.

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