

Predicting Low Risk of Clinically-Important TBI in Children: Vetting and Improving Existing Rules

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Introduction and Problem Motivation

Among children ages 0-14, traumatic brain injuries (TBIs) are a leading cause of injury and death, accounting for over 2100 deaths, 35,000 hospital admissions, and 470,000 emergency department visits annually (Faul, Xu, Wald, & Coronado, 2010). Accurately diagnosing TBIs is thus a crucial problem, especially since the earlier a TBI is identified, the sooner potentially life-saving treatment can be administered.

Typically, TBIs are diagnosed via computed tomography (CT) scans of a patient’s brain. However, performing CT scans carries the risk of radiation-induced malignancy in the patient. This risk is especially pertinent for children, as younger individuals are at a higher risk of developing such malignancies. Therefore, while it is crucial to rapidly and accurately diagnose TBIs, it is also imperative that CT scans are performed only when there is reasonable suspicion the individual has a TBI. Otherwise, the risk of radiation-induced complications from the CT scan will outweigh any potential benefit of having CT scan results.

Nathan Kuppermann and his colleagues explored this problem in their 2009 Lancet paper (“Identification of children at very low risk of clinically-important traumatic brain injuries after head trauma: a prospective cohort study”), seeking to develop interpretable rules that could be used to identify children, who, despite presenting to the emergency room for head trauma, are at low risk of having TBIs and thus can avoid CT scans. The purpose of this present study is to build on their work, developing alternative classification schemes and rule sets to identify children for whom CT scans are unnecessary. With an alternative scheme with high predictive accuracy, one can compare with Kuppermann’s to see if similar rules were isolated, either reinforcing the importance of their identified covariates, or providing a new perspective on what is important in predicting TBI. Additionally, in this analysis, we employ our own data cleaning and preprocessing scheme that we find reasonable, allowing us to assess the stability of their findings subject to perturbations at this stage of the data science life cycle.

In addition to vetting their classification rules, another goal of this analysis is to offer some improvement on their classification strategy. Kuppermann and his colleagues developed rule sets for children aged younger than two and children aged two or older, the reason being that younger children are pre-verbal and thus require a different set of standards to identify TBIs than older children who can verbalize their symptoms (citation). We seek to develop a single age-invariant classifier using covariates common to both primary age groups. We hope that such a classifier will not only identify common predictors of TBI across all ages, but also simplify the decision-making process by obviating the need for a second classifier. Nonetheless, in developing new classifiers, priority will be placed on interpretability, brevity and simplicity, and a low false negative rate.

Data

Data Collection

Before going into extensive detail about the dataset used in this analysis, as well as our cleaning and preprocessing steps, we first offer our own thoughts on what data would be most relevant for vetting previous classifiers and developing an age-invariant classifier.

Besides needing a dataset similar in nature to the dataset on which the original classifiers were based (comprised of children presenting to ER with head trauma, etc.), to vet the previous classifiers, we need a dataset that will enable us to develop classifiers with high predictive accuracy and maximal generalizability so as to compare with previous classifiers. For instance, we would need a dataset containing covariates that are both relevant to the presence of TBIs and can be measured at the time a child is ushered into the ER. From common sense, as well as consulting with Dr. Robert Inglis of UCSF, it seems important information might be the injury mechanism (and a rating of severity), any symptoms such as headache or amnesia, and any results of a physical examination that may indicate neurological impact or skull injury. For both vetting the clinical decision rule and developing an age-invariant rule, it would be preferable to have covariates that are as objective as possible so as to avoid issues with interrater reliability and improve generalizability of the rules to new, real-world settings. It would also be preferable to both have details on covariates invariant to gender, race, ethnicity, or any other such characteristic of the population of interest and a population with gender, age, and racial diversity, to reduce the potential of systematic misdiagnoses in these groups should the rule be applied in future settings. Finally, we wish to identify children who do not need CT scans among a population where there is already some doubt as to whether the CT scan will be worth the risk of radiation induced malignancy. So, the ideal dataset will consist of children who are not critically injured and who could conceivably be TBI-free.

To develop an age-invariant classifier, the above points are certainly relevant. However, specific to an age-invariant classifier, we would also need covariates whose presence or values are not age-dependent.

The dataset used in this analysis is the same one used by Kuppermann and his colleagues to develop their classification rules, consisting of 43399 children who presented to the emergency room for head trauma at one of the PECARN network hospitals between 2004 and 2006. For each child, the data were collected by physicians who filled out standardized sheets detailing the child’s condition, including both questions for the child (‘Do you have a headache?’) and results of a physician examination (‘Palpable skull fracture?’). Additionally, the dataset contains information that would be available after the decision to pursue a CT scan has been made, such as the reasons why a CT scan is being pursued, whether a child will need to be sedated for a CT scan, and whether there were any traumatic findings on the scan. The outcome variable (TBI) was obtained by tracking the child’s progress in the hospital after the initial examination (whether they were admitted for neurological surgery, or died in the ED, for example) or, if the child left the hospital, following-up with the child’s parents, the medical record, emergency department process improvement records, and county morgue records to ensure no TBI diagnoses were missed. In total, there are 125 covariates, including the outcome.

It seems that in this dataset, most of the variables we conjecture are important for predicting TBIs are present, but that there are many covariates whose information would not be known at the time our classifier would be implemented. Additionally, many covariates seem more objective than others (‘Palpable skull fracture?’ vs. ‘Is the child acting normally?’). There are also many variables which are not invariant to age. For instance, the ability to say you have a headache is reserved for verbal children who must be above a certain age. These variables and decisions on what to do with them will be discussed at length in Cleaning and Preprocessing, as well as Post-Processing EDA. The dataset also provides demographic characteristics such as age, gender, race, and ethnicity. This information will be discussed at length in Post-Processing EDA.

Cleaning and Preprocessing

In order to achieve a suitable dataset for the predicting low risk of TBI in children according to the above parameters, we first implement a set of cleaning and preprocessing steps. The overarching goal of our cleaning and preprocessing steps is to keep only covariates relevant to predicting TBI at the time a child would present to the ER, keep only observations for whom this classification rule would be relevant (children for whom it’s a toss-up as to whether they’ll need a CT scan), and to deal with missing values. We present our procedure below, along with relevant judgement calls and their perturbations where applicable.

- Remove columns irrelevant to predicting TBI at the time a child enters the emergency room
 - *EmplType*, *Certification*, *Ind...*, *CT...*, *Finding1*, ... *Finding23*, *EDDisposition*, *Observed*, *AgeTwoPlus*, *AgeInMonth*

- Judgement Call [*injMech*]: drop *InjuryMech*
- * Perturbation: keep *InjuryMech*

This includes *EmplType* and *Certification*, any variable about the reasons why a CT scan was ordered, whether sedation would be needed when performing a CT scan, the traumatic findings from the CT scan, the reason why the child left the ER, or whether the child was observed again to see if they needed a CT scan. We remove the former two variables because domain knowledge says that we should not expect the qualifications of the physician to systematically affect the responses collected. The rest of the variables would either not be available at the time a child first enters the ER, or would be known only after the decision to obtain a CT scan has been made. So, these are not of interest. We also remove an indicator for the individuals being above age 2, as well as the age provided in months, opting to keep the age provided in years. Similarly, according to judgement call [*injMech*], we drop the injury mechanism, a 13-level categorical variable detailing how the injury occurred, opting to keep a severity variable, a 3-tiered categorical variable rating the severity of the injury mechanism as ‘low,’ ‘moderate,’ or ‘high.’

- Remove columns with high levels of missingness
 - *Dizzy*, *Ethnicity*

Dizzy, an indicator of whether the individual felt dizzy, and *Ethnicity*, whether the individual was Hispanic, are both missing in approximately 35% of the observations. *Ethnicity* would be useful for posthoc analysis, but with such a large number of missing values, it may not be particularly informative. *Dizzy* could be a good indicator of TBI, but was deemed too subjective and prone to inter-rater reliability issues according to both Kupperman et al. and Dr. Inglis.

- Remove observations whose *GCS* scores are less than 14

GCS, or Glasgow Coma Score, measures an individual’s level of consciousness on a scale from 3 to 15, and is the sum total of three subscores: an eye score, a verbal score, and a motor score. If GCS is less than 14, the individual is quite injured and would likely need a CT scan anyway. We only keep individuals where the total GCS is 14 or 15.

- Impute and Drop based on GCS scores
 - Judgement Call [*missSubGCS*]: drop observations with total GCS of 14, but missing any of the GCS subcategory scores (*GCSVerbal*, *GCSMotor*, *GCSEye*)
 - * Perturbation: keep the observations
 - Impute a full score into each GCS subcategory when total GCS is 15
 - Judgement Call [*fake15GCS*]: drop observations with total GCS of 15, but suboptimal score in a subcategory
 - * Perturbation: keep the observations
 - Judgement Call [*fake14GCS*]: drop observations with total GCS of 14, but optimal score in every subcategory
 - * Perturbation: keep the observations

GCS score is likely an important predictor of TBI, so it is imperative this variable is as completely filled-out as possible in this dataset. However, it should be noted that there are different standards for obtaining GCS scores for individuals younger than two and those two or older, since individuals younger than two cannot verbalize. We note this as a potential problem to developing an age-invariant classifier.

- Remove observations who were pharmacologically paralyzed, sedated, or intubated at the time of evaluation, or for whom any of this information is missing
 - *Paralyzed*, *Sedated*, *Intubated*

According to domain knowledge, individuals who must be pharmacologically paralyzed, sedated, or intubated at the time of evaluation are likely quite injured and do not represent the population of interest for these classification rules.

- Drop and impute based on variables of altered mental status
 - *AMS*, *AMSAgitated*, *AMSSleep*, *AMSSlow*, *AMSRepeat*, *AMSOth*
 - Judgement Call [*AMS*]: drop observations missing any of the above variables

- * Perturbation: impute missing *AMS* based on presence of any of the subvariables, and drop the subvariables and any remaining missing *AMS* observations

AMS is an indicator representing if the child has altered mental status, and the remaining variables are subvariables indicating the reason why the child has altered mental status (agitated, sleepy, slow to respond, asking repetitive questions, etc.) This is likely an important predictor, so it is important to handle missing values. By judgement call **AMS**, we drop any observation with a missing value in any of these categories, since if the observation is missing it is impossible to truly know a value to impute for *AMS*. As a perturbation, we could impute a 1 in *AMS* if any of the subcategories is affirmative, and 0 otherwise. However, it would be impossible to impute for missing values in the subcategories, so we drop these observations.

- Drop and impute based on variables of other non-head substantial injuries
 - *OSI*, *OSIExtremity*, *OSICut*, *OSICspine*, *OSIFlank*, *OSIAbdomen*, *OSIPelvis*, *OSIOth*
 - Judgement Call **[OSI]**: same procedure as above for *AMS* and its subvariables
 - * Perturbation: same procedure as above for *AMS* and its subvariables

OSI is an indicator on whether the physician determined the child sustained substantial non-head injuries. This could be an injury to the extremities like a fracture, a cut requiring surgery, a spinal cord injury, a flank injury, an intra-abdominal injury, or a pelvis injury. Although not directly related to head trauma, we find this variable might be an indicator of the injury severity. If another variable relevant to head trauma fails to indicate a TBI for a particular individual, including this variable offers another opportunity to diagnose the TBI, possibly suppressing the false negative rate. By judgement call **OSI**, we drop any observations missing any of these values or their subvariables for the same reason as judgement call **AMS**.

- Drop and impute based on hematoma information
 - *Hema*, *HemaSize*, *HemaLoc*
 - Judgement Call **[HEMA]**: same procedure as above for *AMS* and its subvariables
 - * Perturbation 1: same procedure as above for *AMS* and its subvariables
 - * Perturbation 2: Only drop missing *HemaLoc* and *Hema*, impute missing *Hema* from *HemaLoc* presence, drop remaining missing *Hema*

Hema indicates the presence of a scalp hematoma, or swelling due to some head trauma, *HemaSize* is a categorical variable representing the size (< 1cm, 1-3cm, > 3cm) and *HemaLoc* is a categorical variable representing the location (frontal, occipital, parietal/temporal). According to domain knowledge, scalp hematomas are fairly good predictors of TBI. By judgement call **[HEMA]**, we drop all observations missing any of these categories. By Perturbation 1, we only keep the indicators for the presence of hematoma. By Perturbation 2, we keep the indicator as well as the location, since according to domain knowledge, the impact of head trauma to the brain is dependent on the location of the trauma.

- Drop and impute based on skull fracture variables
 - *SFxPalp*, *SFxPalpDepress*
 - Judgement Call **[SFx]**: recode an unclear *SFxPalp* observation as true, drop observations missing either *SFxPalp* or *SFxPalpDepress*
 - * Perturbation 1: same procedure as **[AMS]** perturbation
 - * Perturbation 2: same procedure as **[AMS]** judgement call

SFxPalp is an indicator for a palpable skull fracture, which can also be marked as ‘unclear’ if there is too much swelling to make a determination. *SFxPalpDepress* is an indicator on whether the fracture feels depressed. Both of these are likely important predictors via domain knowledge. As a judgement call, we take an unclear observation as being a palpable skull fracture, since even if the examination was unclear, there is still substantial head trauma.

- Drop missing values for bulging Anterior Fontanelle
 - *FontBulg*

FontBulg is an indicator for whether the child’s anterior fontanelle is bulging, indicating pressure within the brain. This feature would typically only be present in infants, since for older individuals their anterior fontanelle will have closed as their skull hardens with age. While possibly a good predictor for TBI in infants,

this variable does not have much predictive value for older children.

- Drop and impute based on basilar skull fracture variables
 - *SFxBas*, *SFxBasHem*, *SFxBasOto*, *SFxBasPer*, *SFxBasRet*, *SFxBasRhi*
 - Judgement Call [**SFxBas**]: same as [**AMS**] judgement call
 - * Perturbation: same as [**AMS**] perturbation

SFxBas is an indicator if the physician has determined there is basilar skull fracture present, or a fracture at the base of the skull. The rest of the variables indicate common indicators of basilar skull fracture (hemotympanum, otorrhea, raccoon eyes, Battle’s sign, rhinorrhea). More specifically, hemotympanum is when blood is found in the middle ear, otorrhea is when spinal fluid drains from the ear, raccoon eyes is when internal bleeding occurs below the eye, Battle’s sign is a bruise at the bottom of the skull, and rhinorrhea is when spinal fluid leaks out the nose. This is known to be quite an important predictor of TBI, so we wish to make these variables as complete as possible for an ideal dataset. To do so, we follow the same procedure for *AMS* and its subvariables, regarding missingness.

- Drop and impute based on above-the-clavicle injury variables
 - *Clav*, *ClavNeck*, *ClavFro*, *ClavOcc*, *ClavPar*, *ClavTem*
 - Judgement Call [**Clav**]: same as [**AMS**] judgement call
 - * Perturbation: same as [**AMS**] perturbation

Clav is an indicator on whether there was a laceration, abrasion, or hematoma above the clavicles (neck). The subvariables indicate whether there was trauma on the neck or the frontal, occipital, parietal or temporal regions of the scalp. In the same way as *OSI*, this variable will be useful to keep around because it could indicate a TBI if one of the head trauma variables fails to.

- Drop and impute based on neurological deficit variables
 - *NeuroD*, *NeuroDMotor*, *NeuroDSensory*, *NeuroDCranial*, *NeuroDReflex*, *NeuroDOth*
 - Judgement Call [**Neuro**]: same as [**AMS**] judgement call
 - * Perturbation: same as [**AMS**] perturbation

NeuroD is an indicator for whether or not there is some neurological deficit besides altered mental status, as determined by the physician. This includes motor, sensory, cranial nerve (pupil reactivity), or reflex issues. Generally, these categories are quite important for predicting TBI according to domain knowledge. As with previous groups of variables, we implement a judgement call to drop any observation missing any of these values.

- Drop and impute based on vomiting variables
 - *Vomit*, *VomitNbr*, *VomitStart*, *VomitLast*
 - Judgement Call [**Vomit**]: same as [**AMS**] judgement call
 - * Perturbation: same as [**AMS**] judgement call

Vomit is an indicator for whether the individual vomited (as reported by child or parents), *VomitNbr* is the number of times the child vomited post-injury (1, 2, > 2 times), *VomitStart* is how soon after the head injury the vomiting started (within 1 hour, 1 - 4 hours, > 4 hours after), and *VomitLast* is how soon before the evaluation was the last episode (< 1 hour, 1 - 4 hours, > 4 hours before evaluation). As a judgement call, we only keep *Vomit* and impute missing values from the subvariables, since according to domain knowledge, more details about vomiting is generally unhelpful. As a perturbation, we keep these covariates.

- Drop and impute based on headache variables
 - *HA_verb*, *HASStart*, *HASSeverity*
 - Judgement Call [**HA**]: Drop missing *HASSeverity*, impute missing *HA_verb* from *HASSeverity*, and drop remaining missing *HA_verb*, ignore *HASStart* column
 - * Perturbation 1: same as [**AMS**] perturbation
 - * Perturbation 2: same as [**AMS**] judgement call

HA_verb is an indicator on whether the child had a headache, and is marked differently if the child is too young to verbalize that they had a headache, or unable to speak. *HASStart* and *HASSeverity* are both variables indicating when the headache began (before injury, within 1 hour after, 1-4 hours after, > 4 hours after) and

the severity (mild, moderate, and severe). As a judgement call, we keep *HA_verb* and *HASeverity* since these are both more likely to predict TBI than *HASstart*. However, it should be noted that this is an age-dependent variable as discussed in the prior section, since being able to verbalize headache symptoms requires the child to be of a certain age.

- Drop and impute based on seizure variables
 - *Seiz*, *SeizLen*, *SeizOccur*
 - Judgement Call [Seiz]: Drop missing *SeizLen*, impute missing *Seiz* from *SeizLen*, drop missing *Seiz*, and ignore *SeizOccur* column

Seiz is an indicator if the child had a seizure, as reported by the parent or child. *SeizLen* is how long the seizure lasted (< 1 min, 1-5 min, 5-15 min, >15 min) and *SeizOccur* is when did it occur relative to the injury time (immediately on contact, within 30 minutes, or > 30 minutes after injury). Whether a seizure occurred is important to consider, and the length is also somewhat important. So, we opt to keep *SeizLen* and *Seiz*, but drop *SeizOccur*, since this is not as important according to domain knowledge.

- Drop and impute based on loss of consciousness variables
 - *LOCSeparate*, *LocLen*
 - Judgement Call [LOC]: recode ‘suspected’ as ‘yes’ in *LOCSeparate*, and drop observations with missing *LOCSeparate* and *LOClen*
 - * Perturbation 1: same as [AMS] perturbation
 - * Perturbation 2: same as [AMS] judgement call

LOCSeparate is an indicator if the child lost consciousness at all after the injury, and *LocLen* is how long the child lost consciousness (< 5 sec, 5 sec - 1 min, 1 - 5 min, > 5 min). Loss of consciousness is generally a good predictor of head trauma, so we wish to include this to predict TBI. As a judgement call, we consider a suspected loss of consciousness as affirmative, since if there is any doubt as to whether the child lost consciousness, this should be of concern to err on the side of caution.

- Drop missing values for *Amnesia* and *High_impact_InjSev*

Amnesia indicates whether the child has amnesia for the event, but can only be answered by verbal children. This is an example of a variable that is not age-invariant, although it could be a good predictor for verbal children. *High_impact_InjSev* is the rating of the injury mechanism severity, as mentioned above.

- Impute missing values for *ActNorm*
 - Judgement Call [ActNorm]: recode null values as affirmative
 - * Perturbation: recode values as not affirmative

ActNorm is a subjective assessment by the parent as to whether the parent thinks the child is acting normally. Although subjective, domain experts conclude that a parents assesment of their child’s behavior is a good indicator of how severely the child may be injured. Because this is important, we wish to impute as many values as possible. As a judgement call, we consider a missing value as being normal, since if the child was acting normally, it’s possible this question could have simply been skipped over. As a perturbation, we reverse this judgement call.

- Make a pooled outcome variable

We define the outcome, a clinically-important TBI, in the same way as Kuppermann and his colleagues, except we impute missing values in the outcome by the presence of any of the following: death due to TBI, intubated for more than 24 hours, neurosurgery was performed, and the individual was hospitalized for more than 2 nights due to head injury. This should give a complete assessment as to whether an individual had a TBI.

Data Summary

Once we have cleaned the data according to the above steps, we wish to extract binary features from the categorical variables, where applicable. For every subvariable/ parent variable pairing mentioned above (for example, *NeuroDMotor* and *NeuroD*), if the parent variable is marked as false, the subvariable has a value

indicating it is missing. As a default, we maintain this missing category, and binarize the variables by one-hot encoding. Alternatively, we could impute false into the missing values and drop the parent variables, so now each subvariable indicates the presence of the subvariable AND the parent variable as opposed to the subvariable GIVEN the parent variable.

After performing this one-hot encoding, we have 137 features in total. We split the data into a training set (19947 observations), a validation set (6649 observations), and a test set (6649 observations). For post-processing EDA, we only consider the training set, only using the test set to check model accuracy. We should note that there may be cluster structure in the dataset by virtue of the data being collected from different hospitals, presenting a possible comparability issue. However, we are not provided with hospital information, so we perform a random train-validation-test split.

Exploratory Data Analysis

Data Collection detailed a list of characteristics of an ideal dataset to vet and improve upon the original decision rule, according to the goals outlined in Introduction and Problem Motivation. In this section, we explore the relevance of the cleaned and preprocessed data to answer the question at hand, assessing how the data meets these ideals via exploratory plots. The results of our exploratory analysis will then be used to motivate decisions regarding feature inclusion and sample splitting as we build our classifiers.

Of course, the dataset already meets some of these conditions by virtue of our cleaning process: the cleaned dataset only includes individuals with GCS scores of 14 or 15, and individuals who were not so injured as to be intubated, pharmacologically paralyzed, or sedated at the time of evaluation. Additionally, since we are using the same original dataset as Kuppermann et al., where children presented to the ER with head trauma, our results can be suitably compared to theirs. Furthermore, since our data was cleaned differently from theirs, we get an added stability assessment of their decision rule for free, a crucial element of vetting their work.

As mentioned in Data Collection, one of the primary criteria of a good dataset for this problem is a dataset which is diverse in demographic characteristics, including age, race, and gender. This is crucial for generalizability of our work, as a classifier built from a dataset that is demographically lopsided could exhibit unforeseen biases when exposed to more diverse data.

Figure 1 shows distributions of various demographic variables in the cleaned training dataset. In the top-left plot, we see that younger ages are generally represented more than older ages. However, younger children (2 or younger) share many characteristics, including that they are likely preverbal, whereas older children (older than 2) are likely verbal. With this split, it seems like there would be sufficient representation for each age group. Regardless, there are at least 1500 observations for each age out of 19947 total observations, which we find to be sufficient representation.

In the top-right plot, we see that there are about 8000 females and 12000 males in the dataset. While we wouldn't expect different indicators for traumatic brain injury by gender, the dataset is a bit lopsided in this regard; perhaps young boys tend to engage in more dangerous activities than young girls, and thus are at greater risk of head injury. Similarly, in the bottom-right plot we see that the vast majority of observations are White or Black, while other races are quite underrepresented by comparison. Once again, we would not expect a race-dependence of the indicators for TBIs, but like with the observed gender imbalance, we will explore the potential impact of this racial imbalance in Post-Hoc Analysis.

In the bottom left plot, we see that almost 20% of the TBIs observed are in children aged 0, and that the number of TBIs decreases as age increases, flatlining on average for children older than 2. Similarly with the age imbalance, if we were to split by young children versus older children, the number of TBIs in each age group would be more balanced and likely sufficient to generate a classification rule.

While an imbalance in outcome across ages may skew our age-invariant classifier to detect rules relevant for one age group versus another, we should be careful to only include predictors whose relationships with the outcome are invariant by age. The next few plots provide evidence of age dependence and lack thereof for certain variables, motivating the use of certain features for an age-invariant classifier.

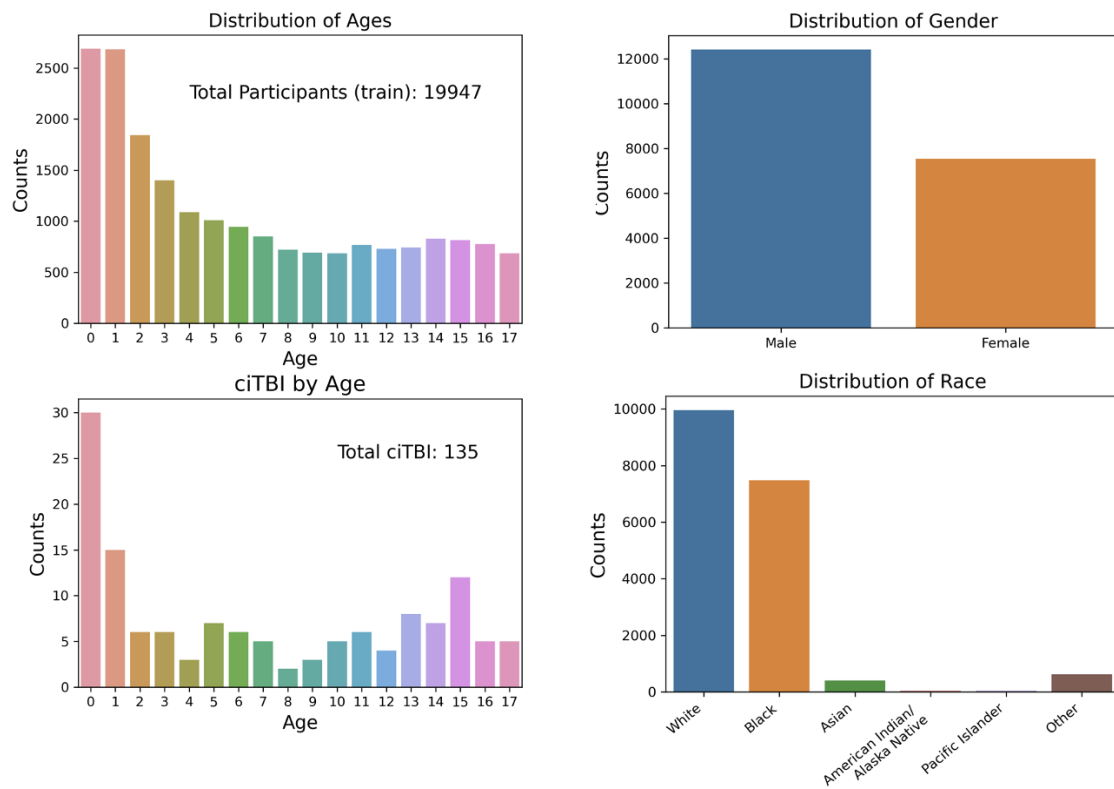


Figure 1: Distributions of various demographic variables in the training dataset, after cleaning and preprocessing.

First, Figure 2 shows the distribution of ‘verbal’ vs. ‘non-verbal’ or ‘pre-verbal’ in response to the physician asking the child if they have amnesia or any headache symptoms. We clearly see that, among children aged two or younger, more than 80% of them were marked as non-verbal in response to both questions. So, a rule based on whether or not the individual has a headache or amnesia would be useless for these individuals. However, among children older than aged two, more than 90% of individuals were able to verbalize answers to these questions. These variables would likely be good predictors of a TBI, so for a classifier only on older individuals, these variables would be useful to include. However, they should not be included in any classifier for younger children, or for an age-invariant classifier.

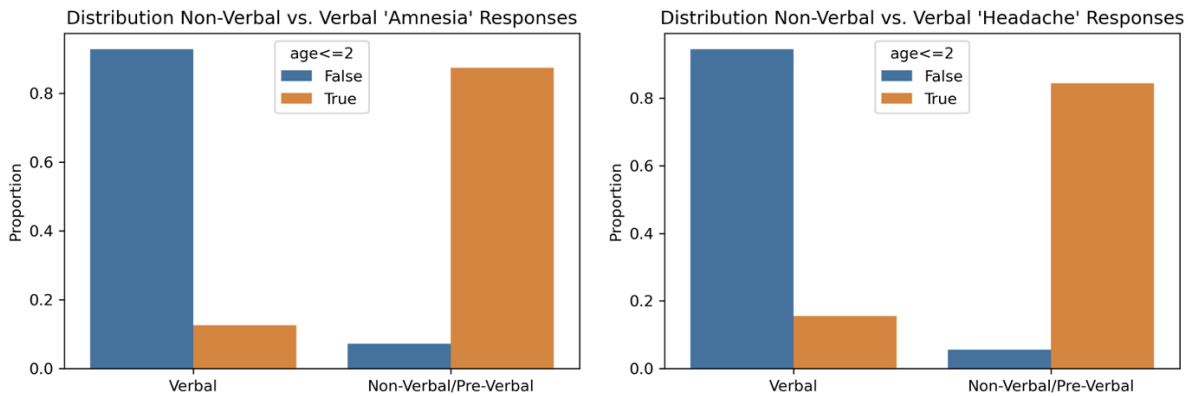


Figure 2: Distributions of ‘Verbal’ or ‘Non-Verbal/Pre-Verbal’ responses to questions on amnesia and headache symptoms, by age.

Next, we recall that, according to Kuppermann et al., for children two or younger, a Pediatric GCS score was assigned, whereas for children older than two, a standard GCS score was assigned. In essence, these are two different standards for evaluating the consciousness of children, since children of different ages have different abilities in expressing their consciousness. However, because the standards are different across these two age groups, we fear that a suboptimal score with regard to one standard might be more indicative of a TBI than a suboptimal score with respect to the other standard. If GCS scores are included in an age-invariant classifier, this presents a potential comparability issue, where the rule that is learned may be applicable to one age-group but not the other. Figure 3 shows the distribution of the outcome variable for suboptimal versus optimal scores of each GCS category (row) for the different age groups (column).

In this figure, we see that, for the GCS Eye scores and the GCS Verbal scores, the outcome conditional on suboptimal versus optimal scores is distributed similarly across both age groups. This indicates that the relationship between GCS score in these categories and the outcome is age invariant. However, for the GCS Motor scores, we see that the distributions are not the same. In fact, it seems like for individuals older than age 2, half of individuals that had a suboptimal GCS Motor score ended up having a TBI, whereas among children two or younger, a much smaller proportion of those with a suboptimal GCS Motor score had a TBI. This tells us that there is likely some age dependence in how this variable is related to TBI. So, when building an age-invariant classifier we also wish to exclude the GCS scores.

Before moving on, we should also establish the age-invariant relationships of other variables of interest with the outcome variable. Figure 4 shows that the distributions of the outcome conditional on having neurological deficits or not, and having basilar skull fracture or not, are not age-dependent. This is as we would expect, since these injuries are observed directly by doctors and are not dependent upon verbal ability or any other factor directly associated with age. Since the relationship of these indicators with the outcome are invariant to age, we should certainly include variables like these in an age-invariant classifier.

Finally, in making a classifier to vet and possibly improve upon that of Kuppermann et al., we explore the potential impact of including and excluding subvariable details. Recall from Cleaning and Preprocessing that

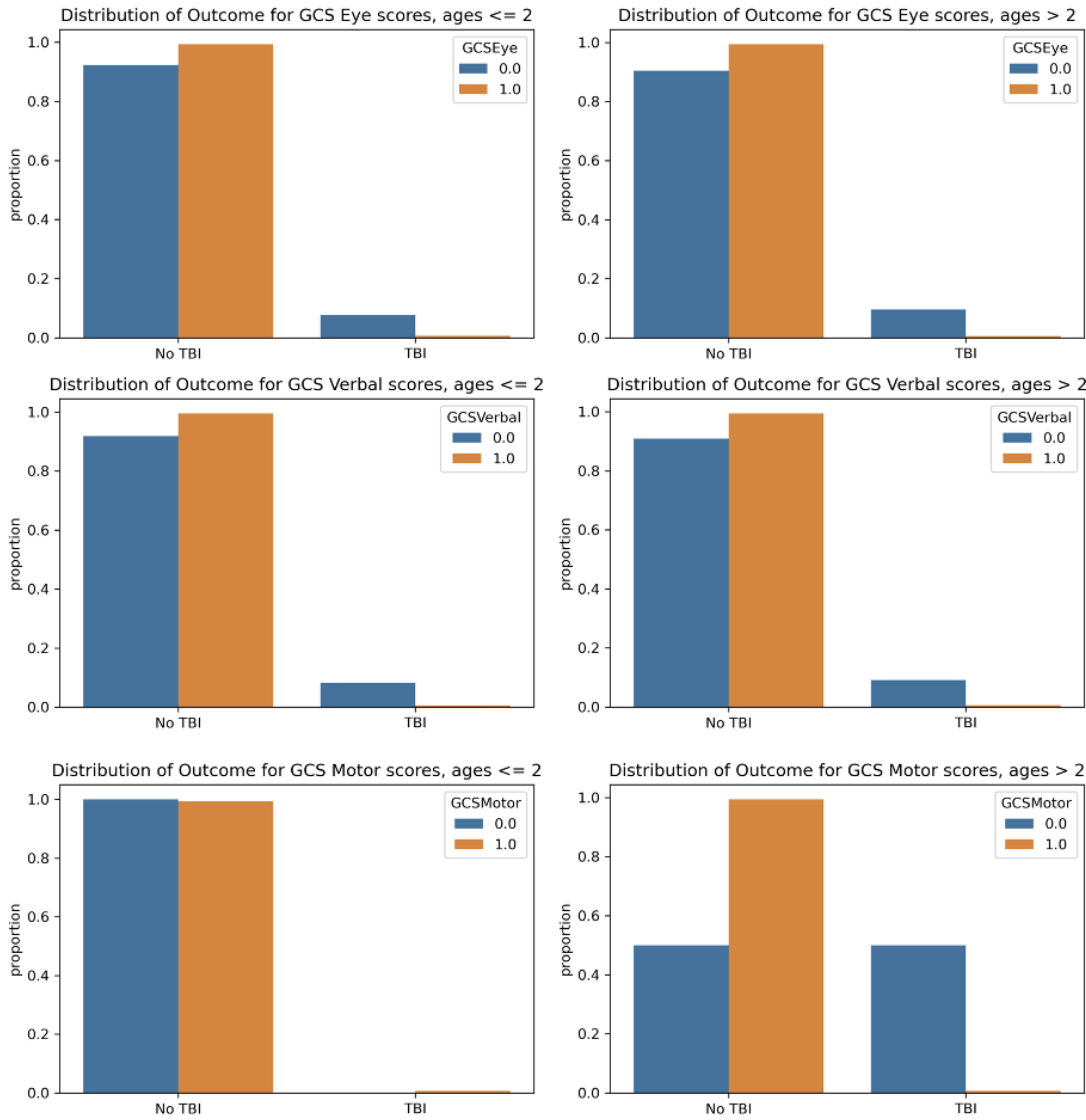


Figure 3: Distributions of outcome among different GCS scores (row) for different age groups (column). 1 indicates optimal score in a GCS category, 0 is one point suboptimal (to achieve total GCS of 14 instead of 15).

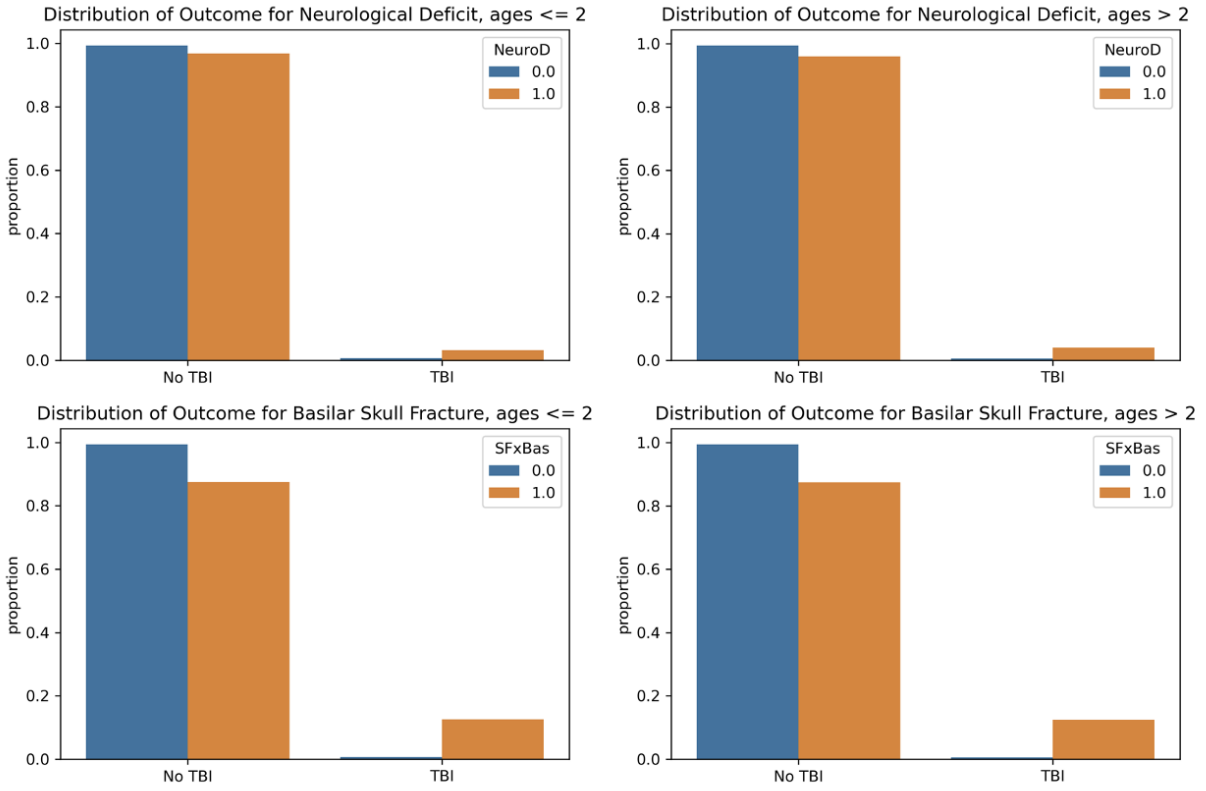


Figure 4: Distributions of outcome conditional on neurological deficit and basilar skull fracture, across both age groups (1 indicates the presence of neurological deficit/basilar skull fracture).

there are many categories which can be thought of as parent categories (Altered mental status? Neurological Deficit? Basilar skull fracture?), with children subcategories that provide more detailed information about the parent category (if altered mental status, is it because you are sleepy? Agitated? Asking repetitive questions?). In making an interpretable classifier with high predictive accuracy and a low false negative rate with which to compare to Kuppermann’s, there are arguments for and against including this subvariable information.

On the one hand, it is possible that certain subvariables are more predictive of the outcome than other subvariables. Consider Figure 5, which shows the correlations of the different subvariables for two parent variables, altered mental status and basilar skull fracture, with the outcome. Each subvariable is binary encoded: if the individual has any of the qualities, it is marked as 1, otherwise, it is marked as 0. For AMS, it seems ‘Sleepy,’ ‘Slow to Respond,’ and ‘Agitated’ are much more correlated with the outcome than ‘Repetitive.’ Even more so, for basilar skull fracture, it seems ‘Hemotympanum’ is much more highly correlated with the outcome than the other variables. If we were to exclude this subvariable information in lieu of only including an indicator for basilar skull fracture, it is possible that we could over-indicate TBIs. Imagine if a child came into the emergency room with CSF Rhinorrhea, but our classifier only considers whether there is a sign of skull fracture. Skull fracture, as an umbrella variable, might be an important predictor, but only because hemotympanum is so correlated with the outcome. Thus, the child would probably be recommended to have a CT scan. However, the child may not have had a TBI in the first place because CSF Rhinorrhea is not very correlated with the outcome. Since we would be giving a child an unneeded CT scan, it could be important to include this subvariable information in an ideal dataset.

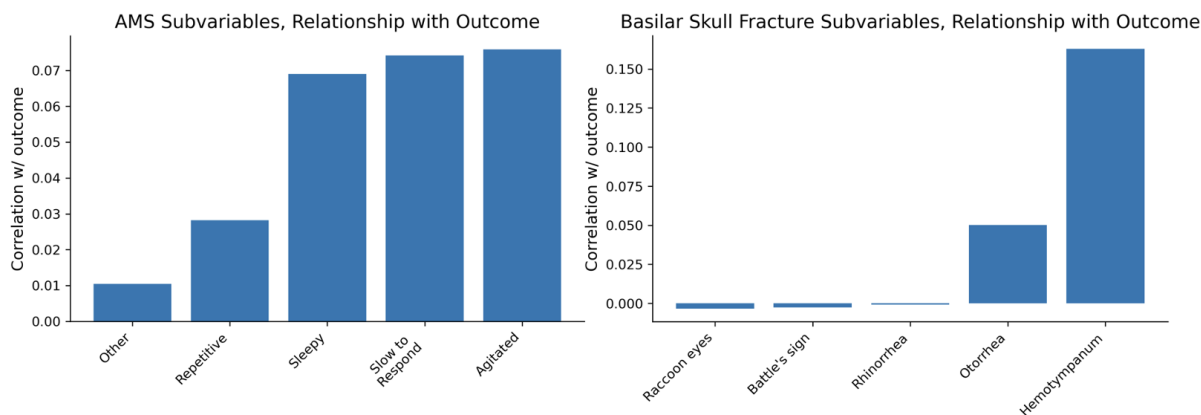


Figure 5: Subvariables of AMS (altered mental status) and basilar skull fracture are correlated differently with the outcome.

However, a clear argument against including this subvariable information is that including too many variables may lead to very uninterpretable and nonsensical rules that are the result of overfitting. Additionally, it seems like the parent variables alone are substantially correlated with the outcome, as is shown in Figures 6 and 7. So, in order to achieve a suitable dataset to make interpretable and accurate classifiers with which to vet and improve upon Kuppermann’s, it seems like it may be suitable to just include the parent variables. Note also that the following two plots further show the dataset is suitable for making an age-invariant classifier: among the age-invariant variables, the correlations with the outcome are almost identical across age groups.

Models

Baseline Model

We choose the model proposed by Kuppermann and his colleagues as the baseline model with which to predict whether a child will have a clinically-important TBI. To evaluate their model, we also use the provided

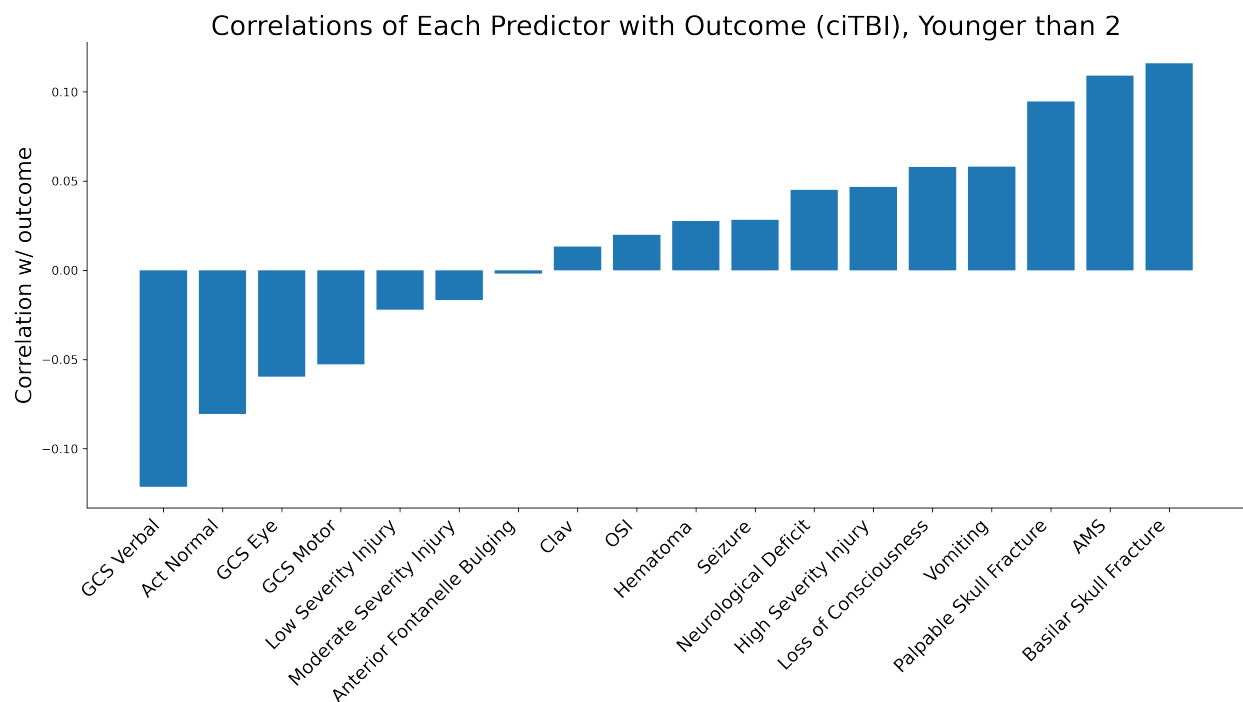


Figure 6: Among children 2 and younger, relevant parent variables seem substantially correlated with the outcome.

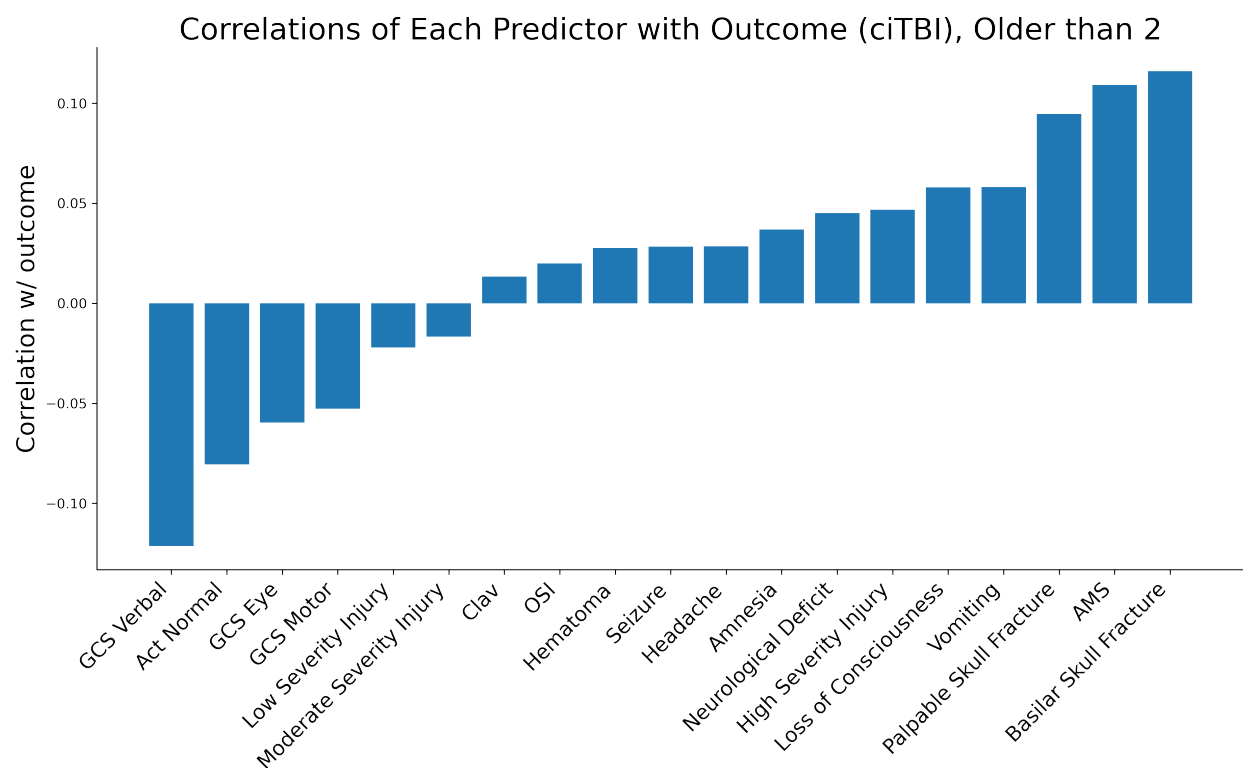


Figure 7: Among children older than 2, relevant parent variables seem substantially correlated with the outcome.

dataset cleaned similarly to theirs, consisting of 42412 children presenting to the emergency room due to the head trauma with GCS scores 14 and above and with the final outcome non-missing. 376 of these observations experienced clinically-important TBIs. In addition, we also use the same age splitting method as Kuppermann, where we use their baseline classifier for preverbal children (< 2 years of age) to predict TBI in this subgroup, and the classifier for verbal children (2 years and older) to predict TBI in this subgroup.

Description

The age-based classifiers in Kuppermann et al. are rule-list models, or a models consisting of a list of rules, ordered by importance in indicating a TBI. A child is labelled as having a TBI if and only if this child satisfies at least one of the rules, where the rules are checked in the aforementioned order of importance. Once we come across a rule that the child satisfies, we stop checking. Therefore, based on the order, each rule is applied to different numbers of children. The list below gives the rules for Kuppermann’s pre-verbal (younger than 2) classifier. Refer to Data Collection for a more in-depth description of the variables mentioned in each rule.

- Altered mental status (*AMS*)? If yes, TBI. If not, continue:
- Occipital, Parietal, or Temporal scalp hematoma (*HemaLoc*)? If yes, TBI. If not, continue:
- Loss of consciousness for more than 5 seconds (*LOClen*)? If yes, TBI. If not, continue:
- Injury mechanism rated severe (*High_impact_InjSev*)? If yes, TBI. If not, continue:
- Palpable or unclear skull fracture (*SFxPalp*)? If yes, TBI. If not, continue:
- Acting normally according to parent (*ActNorm*)? If yes, TBI. If not, no TBI.

The list below now gives the rules for Kuppermann’s verbal (2 or older) classifier.

- Altered mental status (*AMS*)? If yes, TBI. If not, continue:
- Loss of consciousness for more than 5 seconds (*LOClen*)? If yes, TBI. If not, continue:
- Injury mechanism rated severe (*High_impact_InjSev*)? If yes, TBI. If not, continue:
- Basilar skull fracture (*SFxBas*)? If yes, TBI. If not, continue:
- Severe headache (*HA_Severity*)? If yes, TBI. If not, no TBI.

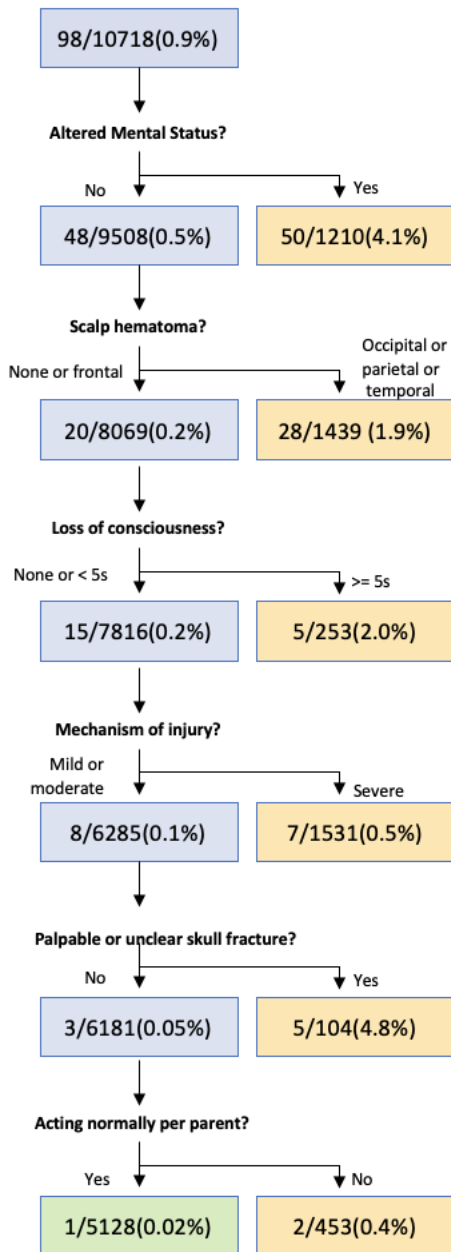
Procedure and Performance

The result of baseline model is shown in 8. The yellow blocks are identified as positive while the green blocks are identified as negative. In each block, the first number represents the number of children experiencing clinically-important TBI, the second number is the number of children in this block, and the last one represents the proportion. And we use the whole cleaned dataset to evaluate the model, while Kuppermann et al. splits the data into training set and validation set. The performance of the baseline classifiers is similar to the result in Kuppermann et al. The baseline classifiers do identify almost all of the cases since the sensitivity is 98.9% and 96.8% for preverbal children and verbal children, respectively. However, the specificity only achieves 48.3% and 57.7%. In our following model, we will propose the one with similar sensitivity and higher specificity. The detailed evaluation metrics based on sensitivity and specificity is discussed in the following part.

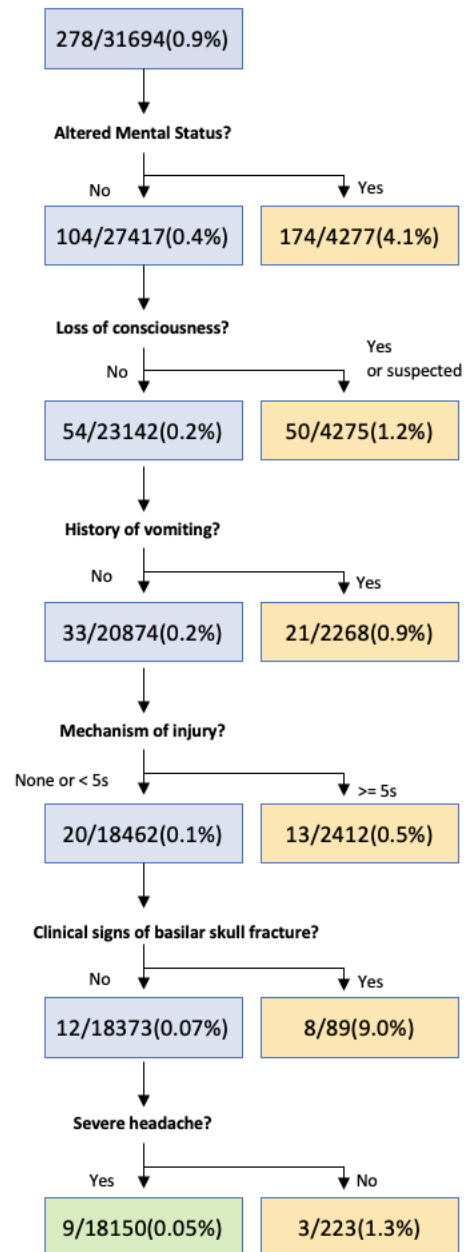
Proposed Model

Describe the proposed model, it’s advantages, disadvantages, how it might compare to the original model, etc.

Justify the proposed model, and refer back to discussion in Post-Processing Discussion for justification while also answering Q8 about any randomness assumptions.



Age < 2



Age >= 2

Figure 8: Decision rule lists for clinically-important traumatic brain injury in children younger than 2 years and in those aged 2 years and older.

Introduction to and an illustration of rulefit

(Andrej)

#TODO: round the numbers, print less rules

Table 1: Basic model rules in order of importance

	rule	type	coef	support	importance
2	Seiz ≤ 0.5 and AMS ≤ 0.5	rule	- 0.8291038	0.8292683	0.3119705
6	Vomit ≤ 0.5	rule	- 1.0290195	0.9297561	0.2629738
9	GCSTotal > 0.5 and SFxPalp ≤ 0.5 and SFxBas ≤ 0.5	rule	- 0.8303451	0.9253659	0.2182148
1	LOCSeparate ≤ 0.5	rule	- 0.4068522	0.7034146	0.1858306
5	Seiz ≤ 0.5 and Hema ≤ 0.5	rule	- 0.2281164	0.6785366	0.1065391
8	GCSTotal > 0.5 and SFxBas ≤ 0.5	rule	- 0.3041677	0.9473171	0.0679510
0	SFxBas ≤ 0.5 and Hema ≤ 0.5	rule	- 0.1385224	0.6785366	0.0646953
4	OSI > 0.5 and High_impact_InjSev_1 ≤ 0.5 and SFxBas ≤ 0.5	rule	- 0.2013573	0.9287805	0.0517873
7	Seiz ≤ 0.5 and SFxBas ≤ 0.5	rule	- 0.2704319	0.9824390	0.0355210
3	High_impact_InjSev_3 ≤ 0.5 and GCSTotal > 0.5 and SFxBas ≤ 0.5	rule	- 0.0116725	0.6926829	0.0053855

Table 2: Maximal model rules in order of importance

	rule	type	coef	support	importance
176	SFxBasRet_0 ≤ 0.5 and NeuroDCranial_1 ≤ 0.5 and OSIPelvis_1 ≤ 0.5 and SeizLen_2 ≤ 0.5 and VomitNbr_2 ≤ 0.5 and AMSAgitated_0 ≤ 0.5 and AMSAgitated_1 ≤ 0.5 and AMSOth_1 ≤ 0.5 and SFxPalpDepress_1 ≤ 0.5	rule	- 1.2058078	0.8758904	0.94975628
1	ActNorm	linear	- 0.7572490	1.0000000	0.2825871
180	SeizLen_2 ≤ 0.5 and SFxPalp_0 > 0.5 and SFxBasHem_1 ≤ 0.5	rule	- 1.2166894	0.9725880	0.80986620
175	ClavPar_1 ≤ 0.5 and SeizLen_3 ≤ 0.5 and AMSSlow_1 ≤ 0.5 and SFxPalpDepress_92 > 0.5	rule	- 0.6275029	0.9000100	0.8882419
181	HemaSize_3 ≤ 0.5 and SFxBasHem_1 ≤ 0.5	rule	- 0.7097578	0.9269378	0.847071
178	SFxBasRet_0 ≤ 0.5 and GCSEye > 0.5 and VomitLast_1 ≤ 0.5	rule	- 0.5118687	0.9340600	0.60270341
34	LOCSeparate_0	linear	- 0.1902846	1.0000000	0.0562658
51	HA_verb_0	linear	- 0.1074405	1.0000000	0.0525110
182	ClavPar_1 ≤ 0.5 and GCSEye > 0.5	rule	- 0.1778378	0.9194906	0.6483862

rule	type	coef	support	importance
177 NeuroDSensory_1 <= 0.5 and HASeverity_2 <= 0.5	rule	- 0.87729333	0.385045	0.1173559
179 GCSVerbal > 0.5	rule	- 0.98413552	0.275252	0.2202880
184 SeizLen_2 <= 0.5 and AMSAgitated_1 <= 0.5 and SFxPalpDepress_1 <= 0.5 and SFxBasHem_1 <= 0.5	rule	- 0.98149162	0.251448	0.1865600
183 SFxBasOto_1 <= 0.5 and AMSOth_0 <= 0.5	rule	- 0.91862721	0.185233	0.0677500
41 LocLen_92	linear	- 1.00000001	0.148932	0.0503670
174 AMSOth_0 <= 0.5	rule	- 0.91878910	0.20802	0.0076152
124 ClavNeck_1	linear	0.00000000	0.00000000	0.00000000
118 HemaSize_3	linear	0.00000000	0.00000000	0.00000000
119 HemaSize_92	linear	0.00000000	0.00000000	0.00000000
120 ClavFace_0	linear	0.00000000	0.00000000	0.00000000
121 ClavFace_1	linear	0.00000000	0.00000000	0.00000000
122 ClavFace_92	linear	0.00000000	0.00000000	0.00000000
123 ClavNeck_0	linear	0.00000000	0.00000000	0.00000000
130 ClavOcc_1	linear	0.00000000	0.00000000	0.00000000
125 ClavNeck_92	linear	0.00000000	0.00000000	0.00000000
126 ClavFro_0	linear	0.00000000	0.00000000	0.00000000
127 ClavFro_1	linear	0.00000000	0.00000000	0.00000000
128 ClavFro_92	linear	0.00000000	0.00000000	0.00000000
129 ClavOcc_0	linear	0.00000000	0.00000000	0.00000000
116 HemaSize_1	linear	0.00000000	0.00000000	0.00000000
131 ClavOcc_92	linear	0.00000000	0.00000000	0.00000000
132 ClavPar_0	linear	0.00000000	0.00000000	0.00000000
117 HemaSize_2	linear	0.00000000	0.00000000	0.00000000
110 SFxBasRhi_1	linear	0.00000000	0.00000000	0.00000000
115 HemaLoc_92	linear	0.00000000	0.00000000	0.00000000
114 HemaLoc_3	linear	0.00000000	0.00000000	0.00000000
97 SFxBasHem_0	linear	0.00000000	0.00000000	0.00000000
98 SFxBasHem_1	linear	0.00000000	0.00000000	0.00000000
99 SFxBasHem_92	linear	0.00000000	0.00000000	0.00000000
100 SFxBasOto_0	linear	0.00000000	0.00000000	0.00000000
101 SFxBasOto_1	linear	0.00000000	0.00000000	0.00000000
102 SFxBasOto_92	linear	0.00000000	0.00000000	0.00000000
103 SFxBasPer_0	linear	0.00000000	0.00000000	0.00000000
104 SFxBasPer_1	linear	0.00000000	0.00000000	0.00000000
105 SFxBasPer_92	linear	0.00000000	0.00000000	0.00000000
106 SFxBasRet_0	linear	0.00000000	0.00000000	0.00000000
107 SFxBasRet_1	linear	0.00000000	0.00000000	0.00000000
108 SFxBasRet_92	linear	0.00000000	0.00000000	0.00000000
109 SFxBasRhi_0	linear	0.00000000	0.00000000	0.00000000
134 ClavPar_92	linear	0.00000000	0.00000000	0.00000000
111 SFxBasRhi_92	linear	0.00000000	0.00000000	0.00000000
112 HemaLoc_1	linear	0.00000000	0.00000000	0.00000000
113 HemaLoc_2	linear	0.00000000	0.00000000	0.00000000
133 ClavPar_1	linear	0.00000000	0.00000000	0.00000000
140 NeuroDMotor_92	linear	0.00000000	0.00000000	0.00000000
135 ClavTem_0	linear	0.00000000	0.00000000	0.00000000

rule	type	coef	support	importance
136 ClavTem_1	linear	0.000000000000000000	0.00000000	0.00000000
157 OSICut_1	linear	0.000000000000000000	0.00000000	0.00000000
158 OSICut_92	linear	0.000000000000000000	0.00000000	0.00000000
159 OSICspine_0	linear	0.000000000000000000	0.00000000	0.00000000
160 OSICspine_1	linear	0.000000000000000000	0.00000000	0.00000000
161 OSICspine_92	linear	0.000000000000000000	0.00000000	0.00000000
162 OSIFlank_0	linear	0.000000000000000000	0.00000000	0.00000000
163 OSIFlank_1	linear	0.000000000000000000	0.00000000	0.00000000
164 OSIFlank_92	linear	0.000000000000000000	0.00000000	0.00000000
165 OSIAbdomen_0	linear	0.000000000000000000	0.00000000	0.00000000
166 OSIAbdomen_1	linear	0.000000000000000000	0.00000000	0.00000000
167 OSIAbdomen_92	linear	0.000000000000000000	0.00000000	0.00000000
168 OSIPelvis_0	linear	0.000000000000000000	0.00000000	0.00000000
169 OSIPelvis_1	linear	0.000000000000000000	0.00000000	0.00000000
170 OSIPelvis_92	linear	0.000000000000000000	0.00000000	0.00000000
171 OSIOth_0	linear	0.000000000000000000	0.00000000	0.00000000
172 OSIOth_1	linear	0.000000000000000000	0.00000000	0.00000000
173 OSIOth_92	linear	0.000000000000000000	0.00000000	0.00000000
156 OSICut_0	linear	0.000000000000000000	0.00000000	0.00000000
155 OSIExtremity_92	linear	0.000000000000000000	0.00000000	0.00000000
154 OSIExtremity_1	linear	0.000000000000000000	0.00000000	0.00000000
144 NeuroDCranial_0	linear	0.000000000000000000	0.00000000	0.00000000
137 ClavTem_92	linear	0.000000000000000000	0.00000000	0.00000000
138 NeuroDMotor_0	linear	0.000000000000000000	0.00000000	0.00000000
139 NeuroDMotor_1	linear	0.000000000000000000	0.00000000	0.00000000
95 SFxPalpDepress_1	linear	0.000000000000000000	0.00000000	0.00000000
141 NeuroDSensory_0	linear	0.000000000000000000	0.00000000	0.00000000
142 NeuroDSensory_1	linear	0.000000000000000000	0.00000000	0.00000000
143 NeuroDSensory_92	linear	0.000000000000000000	0.00000000	0.00000000
145 NeuroDCranial_1	linear	0.000000000000000000	0.00000000	0.00000000
153 OSIExtremity_0	linear	0.000000000000000000	0.00000000	0.00000000
146 NeuroDCranial_92	linear	0.000000000000000000	0.00000000	0.00000000
147 NeuroDReflex_0	linear	0.000000000000000000	0.00000000	0.00000000
148 NeuroDReflex_1	linear	0.000000000000000000	0.00000000	0.00000000
149 NeuroDReflex_92	linear	0.000000000000000000	0.00000000	0.00000000
150 NeuroDOth_0	linear	0.000000000000000000	0.00000000	0.00000000
151 NeuroDOth_1	linear	0.000000000000000000	0.00000000	0.00000000
152 NeuroDOth_92	linear	0.000000000000000000	0.00000000	0.00000000
96 SFxPalpDepress_92	linear	0.000000000000000000	0.00000000	0.00000000
0 Seiz	linear	0.000000000000000000	0.00000000	0.00000000
94 SFxPalpDepress_0	linear	0.000000000000000000	0.00000000	0.00000000
23 InjuryMech_9	linear	0.000000000000000000	0.00000000	0.00000000
25 InjuryMech_11	linear	0.000000000000000000	0.00000000	0.00000000
26 InjuryMech_12	linear	0.000000000000000000	0.00000000	0.00000000
27 InjuryMech_90	linear	0.000000000000000000	0.00000000	0.00000000
28 High_impact_InjSev_1	linear	0.000000000000000000	0.00000000	0.00000000
29 High_impact_InjSev_2	linear	0.000000000000000000	0.00000000	0.00000000
30 High_impact_InjSev_3	linear	0.000000000000000000	0.00000000	0.00000000
31 Amnesia_verb_0	linear	0.000000000000000000	0.00000000	0.00000000
32 Amnesia_verb_1	linear	0.000000000000000000	0.00000000	0.00000000
33 Amnesia_verb_91	linear	0.000000000000000000	0.00000000	0.00000000
35 LOCSeparate_1	linear	0.000000000000000000	0.00000000	0.00000000

	rule	type	coef	support	importance
36	LOCSeparate_2	linear	0.0000000000000000	0.00000000	0.00000000
37	LocLen_1	linear	0.0000000000000000	0.00000000	0.00000000
38	LocLen_2	linear	0.0000000000000000	0.00000000	0.00000000
39	LocLen_3	linear	0.0000000000000000	0.00000000	0.00000000
40	LocLen_4	linear	0.0000000000000000	0.00000000	0.00000000
42	SeizOccur_1	linear	0.0000000000000000	0.00000000	0.00000000
43	SeizOccur_2	linear	0.0000000000000000	0.00000000	0.00000000
44	SeizOccur_3	linear	0.0000000000000000	0.00000000	0.00000000
45	SeizOccur_92	linear	0.0000000000000000	0.00000000	0.00000000
24	InjuryMech_10	linear	0.0000000000000000	0.00000000	0.00000000
22	InjuryMech_8	linear	0.0000000000000000	0.00000000	0.00000000
93	SFxPalp_2	linear	0.0000000000000000	0.00000000	0.00000000
21	InjuryMech_7	linear	0.0000000000000000	0.00000000	0.00000000
2	Vomit	linear	0.0000000000000000	0.00000000	0.00000000
3	GCSEye	linear	0.0000000000000000	0.00000000	0.00000000
4	GCSVerbal	linear	0.0000000000000000	0.00000000	0.00000000
5	GCSMotor	linear	0.0000000000000000	0.00000000	0.00000000
6	AMS	linear	0.0000000000000000	0.00000000	0.00000000
7	FontBulg	linear	0.0000000000000000	0.00000000	0.00000000
8	SFxBas	linear	0.0000000000000000	0.00000000	0.00000000
9	Hema	linear	0.0000000000000000	0.00000000	0.00000000
10	Clav	linear	0.0000000000000000	0.00000000	0.00000000
11	NeuroD	linear	0.0000000000000000	0.00000000	0.00000000
12	OSI	linear	0.0000000000000000	0.00000000	0.00000000
13	Drugs	linear	0.0000000000000000	0.00000000	0.00000000
14	AgeinYears	linear	0.0000000000000000	0.00000000	0.00000000
15	InjuryMech_1	linear	0.0000000000000000	0.00000000	0.00000000
16	InjuryMech_2	linear	0.0000000000000000	0.00000000	0.00000000
17	InjuryMech_3	linear	0.0000000000000000	0.00000000	0.00000000
18	InjuryMech_4	linear	0.0000000000000000	0.00000000	0.00000000
19	InjuryMech_5	linear	0.0000000000000000	0.00000000	0.00000000
20	InjuryMech_6	linear	0.0000000000000000	0.00000000	0.00000000
46	SeizLen_1	linear	0.0000000000000000	0.00000000	0.00000000
47	SeizLen_2	linear	0.0000000000000000	0.00000000	0.00000000
48	SeizLen_3	linear	0.0000000000000000	0.00000000	0.00000000
49	SeizLen_4	linear	0.0000000000000000	0.00000000	0.00000000
73	VomitLast_2	linear	0.0000000000000000	0.00000000	0.00000000
74	VomitLast_3	linear	0.0000000000000000	0.00000000	0.00000000
75	VomitLast_92	linear	0.0000000000000000	0.00000000	0.00000000
76	AMSAgitated_0	linear	0.0000000000000000	0.00000000	0.00000000
77	AMSAgitated_1	linear	0.0000000000000000	0.00000000	0.00000000
78	AMSAgitated_92	linear	0.0000000000000000	0.00000000	0.00000000
79	AMSSleep_0	linear	0.0000000000000000	0.00000000	0.00000000
80	AMSSleep_1	linear	0.0000000000000000	0.00000000	0.00000000
81	AMSSleep_92	linear	0.0000000000000000	0.00000000	0.00000000
82	AMSSlow_0	linear	0.0000000000000000	0.00000000	0.00000000
83	AMSSlow_1	linear	0.0000000000000000	0.00000000	0.00000000
84	AMSSlow_92	linear	0.0000000000000000	0.00000000	0.00000000
85	AMSRepeat_0	linear	0.0000000000000000	0.00000000	0.00000000
86	AMSRepeat_1	linear	0.0000000000000000	0.00000000	0.00000000
87	AMSRepeat_92	linear	0.0000000000000000	0.00000000	0.00000000
88	AMSOth_0	linear	0.0000000000000000	0.00000000	0.00000000

	rule	type	coef	support	importance
89	AMSOth_1	linear	0.0000000000000000	0000000000	0000000000
90	AMSOth_92	linear	0.0000000000000000	0000000000	0000000000
91	SFxFalp_0	linear	0.0000000000000000	0000000000	0000000000
72	VomitLast_1	linear	0.0000000000000000	0000000000	0000000000
71	VomitStart_92	linear	0.0000000000000000	0000000000	0000000000
70	VomitStart_4	linear	0.0000000000000000	0000000000	0000000000
59	HASstart_2	linear	0.0000000000000000	0000000000	0000000000
50	SeizLen_92	linear	0.0000000000000000	0000000000	0000000000
52	HA_verb_1	linear	0.0000000000000000	0000000000	0000000000
53	HA_verb_91	linear	0.0000000000000000	0000000000	0000000000
54	HASseverity_1	linear	0.0000000000000000	0000000000	0000000000
55	HASseverity_2	linear	0.0000000000000000	0000000000	0000000000
56	HASseverity_3	linear	0.0000000000000000	0000000000	0000000000
57	HASseverity_92	linear	0.0000000000000000	0000000000	0000000000
58	HASstart_1	linear	0.0000000000000000	0000000000	0000000000
60	HASstart_3	linear	0.0000000000000000	0000000000	0000000000
69	VomitStart_3	linear	0.0000000000000000	0000000000	0000000000
61	HASstart_4	linear	0.0000000000000000	0000000000	0000000000
62	HASstart_92	linear	0.0000000000000000	0000000000	0000000000
63	VomitNbr_1	linear	0.0000000000000000	0000000000	0000000000
64	VomitNbr_2	linear	0.0000000000000000	0000000000	0000000000
65	VomitNbr_3	linear	0.0000000000000000	0000000000	0000000000
66	VomitNbr_92	linear	0.0000000000000000	0000000000	0000000000
67	VomitStart_1	linear	0.0000000000000000	0000000000	0000000000
68	VomitStart_2	linear	0.0000000000000000	0000000000	0000000000
92	SFxFalp_1	linear	0.0000000000000000	0000000000	0000000000

To see how these relatively basic models perform, we examine their *sensitivity - specificity* curve:

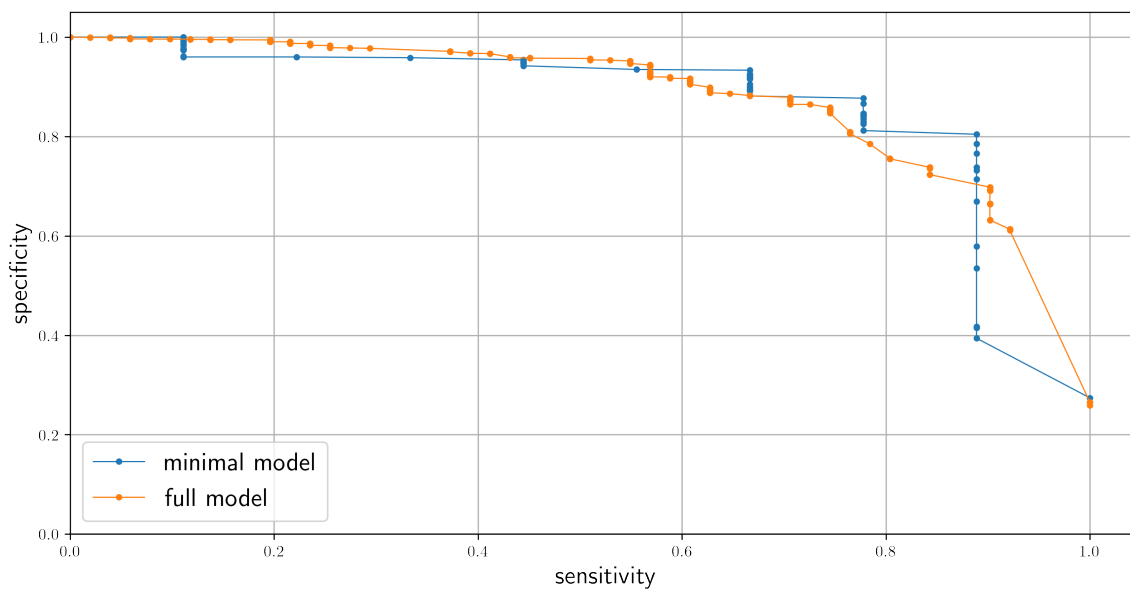


Figure 9: Sensitivity-specificity curve for two basic rulefit models with maximal and minimal judgement calls.

The minimal model is able to achieve 80%+ in both sensitivity and specificity for one threshold value; since

we care much more about false negatives than positives, we consequently value sensitivity far above specificity. Therefore in this case, the fuller model with sensitivities above 90% would be preferred.

////////////////////////////////////

(Xin) I think Andrej will describe rulefit model. Q8 I can only answer this in Logistic regression assumption. Can this be better?

The greedy rule list (CART) model consists of a list of rules, which is similar to the baseline model. Therefore, it is simple to understand, interpret, and pick the important features. Since it is also a greedy method, it does not require distributional assumptions and is computationally-efficient. However, it is non-robust when we perturb the data and the greedy method does not gurantee a good result.

Procedure and Performance

We train and evaluate the models on our cleaned data based on different age-splitting methods and different judgement calls. There are three possible splitting methods in age groups: preverbal children, verbal children, and all children. And we include two different judgement calls: whether include or exclude all subvariables. Although there are six possible datasets, the dataset of preverbal children excluding subvariables contain too few TBI children so we cannot get a reasonable result. Therefore, we will train and evaluate for other five datasets.

For each possible dataset, we split 20% data into the test data, which will only be used in the model evaluation, while we use 80% remained data to fine-tune the hyperparameters and fit the model. Then we split four cross-validation folds to tune the hyperparameters for each model. Since this is an unbalanced dataset, the classification accuracy cannot be a good measure to evaluate the model. The best classifier should identify almost all of TBI cases, as well as identify most of children not experiencing TBI as no TBI. Therefore, we define our evaluation measure as follows.

$$\text{score} = \begin{cases} 5 \times \text{sensitivity} + \text{specificity}, & \text{if specificity} > 0.5. \\ 0, & \text{if specificity} \leq 0.5. \end{cases}$$

For greedy rule list model, the hyperparameters are class weights (representing the importance of children experiencing TBI) and the maximum number of rules in the list (representing the complexity and the interpretability); for rulefit model, the hyperparameter is the maximum length for each rule. In addition, since rulefit model involves random forest methods to pick the rules, the performance may vary even if we choose same hyperparameters. Therefore, for rulefit model, we also need to run the models with different random states and pick the best one as the result, given the maximum length for each rule. Based on the cross-validation results, we pick the best hyperparameters and evaluate the performance on our test data.

The greedy rule list model performs quite bad. For two datasets excluding the subvariables, the largest mean scores in our CV folds are even smaller than 3, which implies the sensitivity is smaller than 60%. For the datasets including subvariables, although the largest mean scores are over 4, the scores on the test data is much smaller. This means the greedy rule list model is very sensitive to the training data and will not behave well on the test data.

The performance of rulefit model on test data is in Figure 10. We can find that the performance for age-invariant group is quite well. As the sensitivity achieves almost 100%, the specificity is larger than 70%. And including subvariables increases the performance of classifying the verbal children. Among two verbal children datasets, when sensitivity is larger than 95%, the specificity is about 70% when we include subvariables, while it drops to 55% if we exclude them. Finally, for preverbal children, the specificity is around 60% when the sensitivity is about 95%. These results imply us developing the age-invariant classifier.

In Figure 10, we lists the important rules for the age-invariant classifier. We find that other signs of altered mental status is the most important linear factor. Besides it, loss of consciousness length and history of vomiting are also two important linear factors. The other important rules consists of several variables. History of vomiting and basilar skull fracture(hemotympanum) forms an important rule. And another important rule

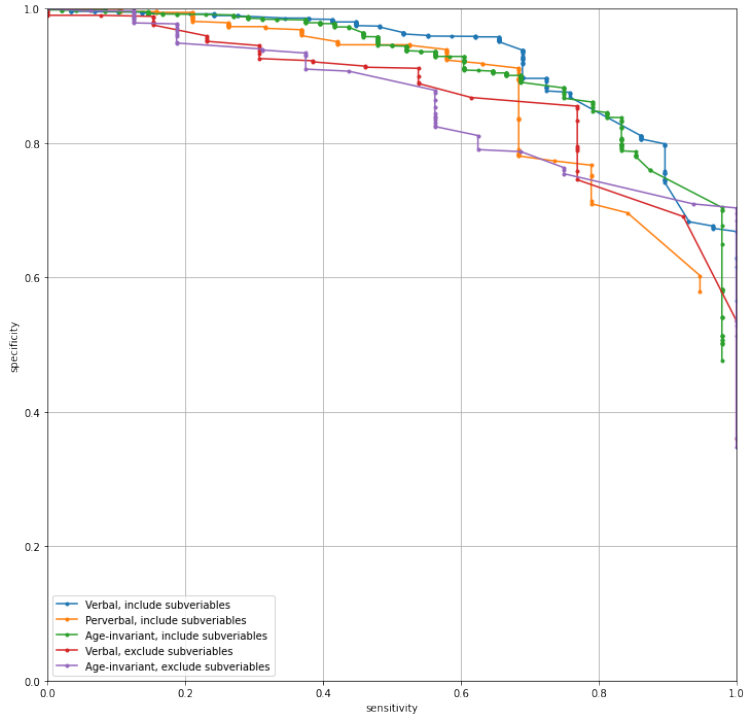


Figure 10: ROC curves of fine-tuned rulefit models on the test data for all five datasets.

	rule	coef	support	importance
127	AMSOth_0 <= 0.5	-0.993656	0.914574	0.277742
124	Vomit <= 0.5 and SFxBasHem_1 <= 0.5	-0.601918	0.873966	0.199769
123	High_impact_InjSev_3 <= 0.5 and LocLen_4 <= 0.5 and SFxPalpDepress_0 <= 0.5 and NeuroDCranial_1 <= 0.5	-0.526590	0.855668	0.185058
121	OSIAbdomen_1 <= 0.5 and HemaLoc_3 <= 0.5	-0.558559	0.897779	0.169209
119	AMSOth_1 <= 0.5 and SFxBasHem_1 <= 0.5 and HemaSize_3 <= 0.5	-0.541056	0.915727	0.150304
18	LocLen_92	-0.391449	1.000000	0.117573
2	ActNorm	-0.296187	1.000000	0.098651
38	AMSOth_92	-0.219569	1.000000	0.065860
125	SFxPalpDepress_92 > 0.5 and SFxBasPer_0 <= 0.5	-0.641207	0.992781	0.054283
129	OSIOth_1 <= 0.5 and SeizLen_2 <= 0.5 and SFxPalpDepress_92 > 0.5 and SFxBasHem_1 <= 0.5 and SFxBasOto_1 <= 0.5	-0.230229	0.974683	0.036166
131	SFxPalpDepress_1 <= 0.5 and SFxBasHem_1 <= 0.5 and NeuroDCranial_1 <= 0.5	-0.314930	0.993483	0.025341
130	ClavPar_1 <= 0.5	-0.061043	0.926054	0.015974
126	OSICut_1 <= 0.5 and SFxPalp <= 0.5	-0.101669	0.978242	0.014833
128	OSICut_1 <= 0.5 and LocLen_4 <= 0.5 and SeizLen_4 <= 0.5 and SFxBasHem_1 <= 0.5 and NeuroDMotor_1 <= 0.5	-0.120065	0.989372	0.012312
122	SeizLen_1 <= 0.5 and SFxPalp <= 0.5	-0.075998	0.973129	0.012289
120	AMSSlow_1 <= 0.5 and SFxPalpDepress_92 > 0.5	-0.043991	0.976989	0.006596

Figure 11: The important rules for the age-invariant classifier

involves the severity of injury, loss of consciousness length, feeling depressed in palpable skull fracture, and Neurological deficit(cranial nerve).

Stability Analysis

Answer Question 9

Go through the perturbations of our judgement calls (which will have been introduced above) and compare the performance of our classifier, see if it is similar or different

Also, if there was instability in training the classifiers (as Xin noted in his notebook in our 12/2 meeting), run through it here as well

Discussion

Discuss the performance, the results, the stability subject to perturbations, how well it compared to the baseline, etc.

Discuss future directions on what we can do (other models, other perturbations, things we missed that we would have done with more time, etc.)

Conclusion

Just wrap it up