**Identifying patients who will not re-achieve remission after breakthrough seizures**

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**Summary**

**Objective:** We aim to identify people with epilepsy who are unlikely to re-achieve a 12 month remission within 2 years after experiencing a breakthrough seizure following an initial 12 month remission.

**Methods:** We apply a novel longitudinal discriminant approach to data from the SANAD study to dynamically predict the risk of a patient not achieving a second remission after a breakthrough seizure by combining both baseline covariates (collected at the time of breakthrough seizure) and follow-up data.

**Results:** The model classifies 83% of patients. Of these, 73% of patients (95% CI: 58%­−88%) who did not achieve a second remission were correctly identified (sensitivity), whilst 84% of patients (95% CI: 69%−96%) who achieved a second remission were correctly identified (specificity). The area under curve (AUC) from our model was 87% (95% CI: 80%−94%). Patients who did not achieve a second remission were correctly identified on average after 10 months of observation post-breakthrough.

Occurrence of seizures after breakthrough and the number of seizures experienced were the most informative longitudinal variables. These longitudinal profiles were influenced by the following baseline covariates: Age at Breakthrough seizure, presence of neurological insult and number of AED’s required to achieve first remission.

**Significance:** Using longitudinal data gathered during patient follow up allows more accurate predictions than using baseline covariates in a standard Cox model. The model developed in this paper is a useful first step in developing a tool for identifying patients who develop drug-resistance after an initial remission.

Key Points

* The presence and number of post-breakthrough seizures are the most informative variables for predicting poor outcome following first breakthrough.
* Age at breakthrough seizure, neurological insult and numbers of treatments required to achieve first 12-month remission are also influential.
* Real time monitoring of seizure history leads to more accurate predictions than those estimated from existing models at baseline, with approximately 10 months of observation required on average to correctly identify patients who do not re-achieve remission.

**INTRODUCTION**

Approximately 30-40% of patients with epilepsy will never enter a sustained remission from seizures despite multiple treatment changes.1 Of the 60-70% of patients who achieve a 12-month remission from seizures, around 37% may have a breakthrough seizure, defined here as one that occurs following a remission of at least 12 months, despite continued treatment with antiepileptic drugs (AEDs).2

Following a breakthrough seizure patients may go on to achieve a further period of remission, either immediately or following changes to their treatment regimen.3 However, some continue to experience seizures despite multiple treatment changes.

If we were able to reliably predict prognosis following a breakthrough seizure, patients could be provided with important information that could influence life choices, and medical resources could be used more effectively, for example patients could be put on a pathway for respective surgery earlier in their disease course.

However, very little work has been conducted to investigate patient outcomes following breakthrough seizures. Bonnett et al4, 5 consider prognostic factors for the risk of a breakthrough seizure following a period of remission, the risk of seizure recurrence following a breakthrough seizure and the chance of achieving a 12-month period of remission following a breakthrough seizure. These studies consider patient data up until the breakthrough seizure as prognostic variables, but not data collected during subsequent follow up. In addition, one study considered prognostic factors that affect seizure relapse and the development of drug resistance in patients who had experienced long term remission.6 Two further studies investigate breakthrough seizures in Uganda7 and Egypt,8 although neither study considered outcomes following the breakthrough seizure.

A few studies have attempted to predict patient’s long-term epilepsy status with various length of observation.9-11 Hughes et al12 describe a model which identifies patients who will not achieve a 12-month continuous seizure free period within 5 years of initial diagnosis. However, this study only considered patients up until their first remission if a remission was observed. Keller et al13 identifies patients who will continue to experience seizures following brain surgery. To the best of our knowledge, no models exist that aim to dynamically identify patients who will not achieve remission following a breakthrough seizure.

Our aim in this study was to describe in a statistical model the seizure history of patients following a breakthrough seizure and use the model to give quantitative predictions. Therefore, further evidence could be added to support clinician intuition regarding the trajectory of patients not likely to achieve a second remission after a breakthrough seizure.

**METHODS**

The Standard and New Antiepileptic Drugs trial (SANAD) has been described in detail elsewhere.14, 15 In summary, the trial was designed to compare two standard antiepileptic drugs with a range of alternatives. Patients were eligible for the trial if they were at least five years old and had experienced at least two clinically definite unprovoked seizures in the last year. Patients for whom carbamazepine was considered to be the standard optimal treatment were recruited to Arm A of the SANAD trial and were randomly allocated in equal proportions to receive either carbamazepine, gabapentin, lamotrigine or topiramate. From 1st June 2001, a further drug, oxcarbazepine, was added to the trial and patients were allocated to these five drugs with equal proportions. Patients for whom valproate was considered the standard optimal treatment were included in Arm B of the trial and were randomly assigned with equal proportions to valproate, lamotrigine or topiramate.

The SANAD study has previously been used to investigate time to treatment failure from randomisation and time to 12-month remission from randomisation.16-18

SANAD is the largest prospective study in patients with epilepsy to date and contains follow-up data for up to 7 years, allowing an excellent opportunity to investigate time-dependent factors that influence the risk of having drug resistant epilepsy.

This analysis describes a model that combines both baseline covariates (recorded at the time of breakthrough seizure) and subsequent follow-up data in order to identify patients who will not achieve a second period of 12-month remission within 2 years following a breakthrough seizure. A patient’s classification is determined by the likelihood that the patient will achieve a second remission or not, and the estimation of this probability can be updated with new data while the patient remains under observation and is not classified as achieving a second remission.

This analysis considers both arms of the SANAD study simultaneously. Since the allocation to arms A or B was dependent on the type of epilepsy, we consider type of epilepsy as a potential baseline covariate. Patients who experienced a 12-month continuous seizure-free period since randomisation, and then subsequently had a breakthrough seizure were eligible for inclusion in this analysis.

**Statistical Analysis**

The primary outcome of interest is whether or not a patient experiences a further 12-month remission within two years of experiencing a breakthrough seizure.

We considered four variables that were recorded at each follow-up visit and modelled the changes in these variables over time using a multivariate generalised linear mixed model (MGLMM).19 Separate models were fitted to patients who are known to have achieved another remission period of 12 months within two years following breakthrough and those who did not. Specifically we consider (1) whether a patient had experienced seizures since their previous clinic visit (yes/no), (2) how many seizures were experienced since the previous clinic visit, (3) the number of patient reported adverse events experienced since the previous clinic visit and (4) whether a patient’s treatment was changed at the last clinic visit (yes/no). Treatment change included changes in dose (increase or decrease) of a drug or the addition or removal of a drug. Potential adverse events included depression, dizziness, allergic reactions, headaches and tiredness amongst others.20

Each variable that was measured at repeated clinic visits (longitudinal variable) was allowed to depend upon the following baseline variables: Age at breakthrough seizure, sex, electroencephalogram (EEG) result at randomisation, computerised tomography (CT) or magnetic resonance imaging (MRI) result at randomisation, type of epilepsy, first degree relative with epilepsy, neurological insult (learning difficulties or a neurological deficit), total number of seizures prior to first remission, total number of treatments prior to first remission, time to first period of 12 month remission, number of tonic-clonic seizures prior to breakthrough seizure, and post breakthrough seizure treatment decision.4, 5 In addition, time since the last clinic visit was included to account for the fact that clinic visits were not equally spaced. Baseline covariates from the above list were included in a forward selection approach with models compared using penalised expected deviance (PED).21 The best combination of baseline covariates to explain each longitudinal variable was determined.

These two fitted models were subsequently used in a longitudinal discriminant analysis22 to classify patients as either likely to achieve another period of 12-month remission within 2 years of breakthrough seizure or likely to not re-achieve 12-month remission within 2 years. A longitudinal discriminant analysis assesses the likelihood that the data of a new patient was generated by each of the two multivariate mixed models. In this sense, the model assesses in a probabilistic manner which of the two average group trends the new patient is closest to.

At each follow up visit for a patient, between their initial breakthrough seizure and their two-year post breakthrough end point, their risk of not achieving remission within two-years of breakthrough seizure was updated using the additional information collected at the visit. The best combination of the four longitudinal variables to be used to classify new patients was selected using the probability of correct classification (PCC), in order to maximise classification accuracy.

Credible intervals around the calculated probability of not achieving remission within 2 years of breakthrough were used to assess the precision of the estimated probability. At each follow up visit, if the calculated credible interval was entirely above a threshold of 0.38 (threshold determined following a ROC curve analysis as the point on the ROC curve closest to the top-left corner of the plot) then the patient was classified as not going to achieve remission within the remaining period of the two years since their breakthrough seizure. If this was not the case then the patient simply remained under observation. At the final follow up visit before 2-year status was confirmed, there was an additional option of classifying a patient as likely to achieve remission if the credible interval was entirely below 0.38. If the credible interval contained the threshold at this visit, then the patient remained unclassified as there was insufficient confidence to predict the patient’s status. Figure 1 gives a diagrammatic representation of this classification scheme.

FIGURE 1 ABOUT HERE

Patients were predicted as likely to not achieve remission, only at the point in follow up at which their credible interval for risk of drug resistance is entirely above 0.38. This means that not all patients are classified at the same time and classification only occurs when there is reasonable confidence that a patient will truly not achieve remission within 2 years of breakthrough.

To test the predictive accuracy of our model, data from 70% of the patients in each group (those who achieved a second remission and those who did not) was used to train the model, and the remaining 30% was used to test the predictive accuracy. This was repeated for 100 random splits of the data into training and test sets. Predictive accuracy measures were then calculated and averaged.

For comparison purposes we compared our longitudinal model with a Cox proportional hazards model described in Bonnett et al.23 The Cox Model predicts at baseline (time of breakthrough seizure) the probability of not achieving remission within 2 years.

**RESULTS**

Out of 2437 patients who were considered for this analysis, 1901 patients were excluded for a number of reasons; in 58 patients the seizures were later linked to causes unrelated to epilepsy (3%), 786 patients did not achieve remission during the follow up period (41%) and 1057 patients did not experience a breakthrough seizure (56%). In total 536 patients experienced a breakthrough seizure (34% of all patients who experienced remission). Patients who had a dose decreased prior to a breakthrough seizure were also excluded from this analysis (n=26, 4.9% of all patients who experienced a breakthrough seizure) since their seizure could potentially be due to antiepileptic drug withdrawal4, 5. A further 210 patients were excluded because, although they did experience a breakthrough seizure, they were not followed for sufficient time to determine their two year status post-breakthrough. Of the remaining 300 patients who experienced a breakthrough seizure, 185 patients (62%) went on to achieve a further period of 12 month seizure remission within 2 years of experiencing their breakthrough seizure and 115 patients (38%) were observed for 2 years following breakthrough seizure without experiencing a 12 month remission.

Table 1 describes the characteristics of patients who were observed to achieve a second period of 12-month remission within 2 years of breakthrough and those who did not. The best combination of longitudinal variables to achieve optimal classification accuracy was a bivariate model including whether the patient experienced seizures or not since their last visit and the total number of seizures experienced since the patient’s last visit. Separate bivariate models were fit to the patients who were observed to achieve remission and those who did not. The two bivariate models were used in the longitudinal discriminant analysis.

TABLE 1 ABOUT HERE

The covariates used to model each longitudinal variable are shown in Table 2. For both groups of patients (those who achieved remission and those who did not), the likelihood of experiencing seizures decreased (odds ratios less than 1) as time since breakthrough increased. Conversely, as the time since last follow up increased the likelihood of experiencing seizures increased (odds ratios greater than 1), probably reflecting the fact that they had a longer period in which to experience a seizure. Time since breakthrough and since last follow up were also associated with the expected number of seizures but with a minor effect. Patients who ultimately achieved second remission, but had required more antiepileptic drugs (AEDs) to achieve their first period of 12 months remission, were expected to experience slightly more seizures than similar patients requiring fewer AEDs (PE=0.207, 95%CI: 0.116-0.302, implying that for each drug required to achieve a patient’s first 12-month remission, the number of seizures they were expected to have experienced since their last, post breakthrough follow up visit increased by 0.207). Patients who had neurological insult on diagnosis, but ultimately achieved remission, were approximately four times more likely to experience seizures than those who did not have neurological insult. For patients who would not achieve a second remission, increasing age increased the risk of experiencing seizures, although the result was not statistically significant.

TABLE 2 ABOUT HERE

To demonstrate how our model works we describe the clinical follow up post breakthrough of three patients (Figure 2). Patient (a) was correctly identified to achieve a second remission. At their first visit, 39 days post breakthrough they reported having experienced 2 seizures, giving a probability of not achieving remission of 0.23 at this visit. However, at two subsequent visits they reported no seizures, and accordingly the model gives a low probability that they will not achieve a second remission. This is confirmed at their next visit, when it is observed that they have been seizure free for 12 months.

FIGURE 2 ABOUT HERE

Patient (b) was correctly identified by the model as not likely to achieve a second period of 12-month remission following a breakthrough seizure. At their first two visits post breakthrough (81 and 165 days), they described experiencing 2 and 10 seizures since the previous visit, respectively. At the second visit the model assigns a high probability of not achieving a second remission to this patient. In addition, the credible interval around this probability is entirely above the threshold of 0.38, and the patient is classified as not going to achieve remission within two years of breakthrough. This prediction was made at 165 days post breakthrough.

Patient (c) was unclassified by our model despite ultimately being observed to achieve remission, since the credible intervals were too wide to determine with confidence the patient’s status (grey shaded area).

The predictive accuracy of our model is assessed by considering how many of the patients were correctly classified (Table 3). Out of the patients classified by the model, 73% of patients (95% CI: 58%-88%) who would not achieve a second remission were correctly identified (sensitivity), whilst 84% of patients (95% CI: 69%-96%) who achieved a second remission were correctly identified (specificity). Overall, 80% of patients (95% CI: 71%-89%) were correctly identified (PCC). The area under curve (AUC) from our model was 87% (95% CI: 80%-94%) showing that the model achieves a good level of discrimination. Out of the patients predicted not to achieve remission, 73% of patients (95% CI: 57%-91%) were observed not to achieve remission (Positive predictive value, PPV), whilst 85% (95% CI: 77%-93%) of patients predicted to achieve remission went on to achieve remission (negative predictive value, NPV). The prediction times reported in Table 3 show the average time at which a patient is correctly identified as not going to achieve remission from seizures, and show that our model is able to identify patients who will not achieve a second remission approximately 10 months after a breakthrough on average.

TABLE 3 ABOUT HERE

The predictive accuracies reported above are based on patients who were classified by the model. Approximately 17% of patients (95% CI: 16%-18%) were left unclassified by our model as there was considerable uncertainty about their status and longer follow up would have been required. If unclassified patients were considered as incorrectly classified, the predictive accuracy would indicate a sensitivity of 57% (95% CI: 41%-71%), specificity of 72% (95% CI: 59%-87%) and PCC of 66% (95% CI: 57%-74%). A key point of this approach is that by leaving a relatively small proportion of patients unclassified, much greater predictive accuracy is obtained for patients who are classified.

For comparison purposes, we compared our model predictions to predictions from the Cox proportional hazards model described in Bonnett et al.23 The predictive accuracy of the Cox model (used to predict chance of experiencing remission within 2 years) is also shown in Table 3, and the corresponding ROC curves are shown in Figure 3. The longitudinal model achieves substantially better classification accuracy. This is emphasised by the boxplots in the left panel of Figure 3, which show much greater separation in the probabilities assigned to patients in each group for the discriminant model than the Cox proportional hazards model. This demonstrates that the information available at the point of breakthrough seizure is insufficient to determine whether or not a patient will go on to achieve remission again. The additional information collected during follow up and incorporated into our longitudinal discriminant analysis model enables more accurate predictions of long-term outcome to be made than by simply using baseline information.

FIGURE 3 ABOUT HERE

**DISCUSSION**

We have shown that the longitudinal information collected during follow-up following a breakthrough seizure, in addition to baseline variables, can be used to identify patients who will not achieve a second period of 12-month remission. This will be intuitive to clinicians who observe patients during follow up and recognise the trajectory of patients likely or unlikely to achieve a seizure remission. In this respect, our model provides further quantitative evidence alongside clinical intuition to support decision making.

We have identified that whether a patient has seizures and the number they have between clinic visits are useful indicators of whether or not they will ultimately achieve a second period of 12-month remission. In addition a patient’s age at the breakthrough seizure, the number of treatments required to achieve their first period of 12-month remission, and whether or not a patient has a neurological insult all have an impact on the likelihood and frequency of seizures experienced following a breakthrough seizure, even if a patient will ultimately achieve a second period of 12-month remission. In this analysis we only considered the number of treatments required, rather than the specific treatments since the number of treatment combinations after withdrawal of randomised drug is too large to model accurately, and differences between the treatments can be small.

Of the patients classified, our model correctly identifies 73% of patients who will not achieve a second period of remission and 84% of those who will. On average our model classifies those not likely to achieve a second remission after approximately 10 months follow-up. Compared to a Cox model, which uses only data up to the time of the breakthrough seizure, our new model is considerably more accurate. This suggests that there is insufficient evidence to predict with accuracy which patients will re-achieve remission on the day of the breakthrough seizure and only by observing patients over time can increased confidence be gained. Although this necessitates a delay in being confident about which patients will not achieve remission, we observed that only approximately 10 months of further observation were required.

**LIMITATIONS**

Since we required patients to have experienced a 12-month remission, and a breakthrough seizure with an additional two years of follow up post seizure the sample size for our analysis is relatively small. These factors potentially limit the power of our analysis. With larger patient groups, additional baseline covariates relevant in the longitudinal evolution of a patient’s seizure history may have been identified. In addition, further longitudinal makers could have been seen to be important predictors of whether or not a patient will re-achieve remission following a breakthrough seizure.

Hughes et al12 used the SANAD data to develop a model for identifying patients who will not achieve a 12-month remission within five years of an initial diagnosis. The reason for the shorter time frame in this analysis was due to the small number of patients with sufficient follow-up observations following a breakthrough seizure to consider a longer period (the median follow up time post breakthrough for the 536 patients who experienced a breakthrough seizure was 1.6 years, Inter quartile range = 0.79-2.57 years)

With longer follow up a longer time frame than 2 years could have been considered. Many of the patients that did not achieve remission within 2 years, may have done so if they had been observed for longer. With a longer period of follow up better predictive accuracy may be achieved.

There is a potential bias in our findings since patients with early first remissions are more likely to be included than patients who achieved a first remission after a longer follow up, due to the need for a further two years of observation in order for the patient to be included in our analysis. Similarly, patients who experience a breakthrough seizure closer to their initial remission are more likely to be included than patients who remained in an initial state of remission for longer. Our analysis omitted 210 patients who experienced a breakthrough seizure but were not observed for a further two years largely due to the end of the SANAD trial. Since we could not determine the status of these patients, we could not include them in the analysis. Studies with much longer follow up would be able to assess this bias.

We excluded patients whose AED dose had been decreased before a breakthrough seizure was experienced. This was because the breakthrough seizure may have been due to drug withdrawal. The predictive tool presented in this paper may not be applicable then to people who have experienced breakthrough seizures following a decrease in AED dose.

For the model presented in this paper to be useful in clinical practice, external validation is required. However, there are no datasets available with the relevant information required to validate our model currently. Internal validation of our model suggests good classification performance.

The SANAD data relies on patient reported seizure counts. It may be the case that these are underreported or, in the case of experiencing many seizures, approximated. This may result in our estimates being an underestimation of the actual numbers of seizures experienced which may in turn bias downwards the estimates of the evolution of total seizure counts over time. If patients who do not achieve a second 12-month remission are expected to experience more seizures, then more accurate seizure counts would increase the separation between the two groups and lead to more accurate classification.

**CONCLUSION**

Hughes et al12 developed a longitudinal discriminant analysis model to predict patients who will not achieve a first period of 12-month remission within five years of initial diagnosis (sensitivity=95%, specificity=97%, AUC=95% for classified patients.). In this paper we go a step further by developing a discriminant analysis model to identify patients who, having initially achieved 12-month remission and gone on to have a breakthrough seizure, will not achieve another period of 12-month remission within 2 years of the breakthrough seizure. We believe that the prediction accuracy of this model is reasonably good, suggesting the potential of this approach to identify patients who will develop drug resistance following an initial remission, although a larger patient sample, and longer follow up would be required to explore this further.

The predictive model here developed, and the one of Hughes et al12 provide a useful tool to identify patients who are not likely to achieve remission from seizures early on in their clinical follow up, both after diagnosis and after a breakthrough seizure following remission. Incorporating these models into an easy to use calculator (possibly as a webtool or App) would be a necessary next step in making these models clinically useful Such an approach has the potential to provide clinicians with more accurate information regarding the long-term outcome of their patients, which could lead to improved patient counselling and more informed treatment decisions, including the possibility of aiming for a most tolerated AED regimen rather than total seizure freedom if this was deemed to be unlikely.

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**Disclosure of Interests:**

None of the authors has any conflict of interest to disclose.

**Ethical Publication Statement:**

We confirm that we have read the Journal’s position on issues involved in ethical publication and affirm that this report is consistent with those guidelines.

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Figure Legends:

Figure 1: Allocation Scheme for identifying patients who will not re-achieve remission within 2 years of breakthrough seizure.

Figure 2: Three individual patient’s probabilities of not achieving a second remission. The crosses show the probability assigned by the model at the clinical visits and the grey shaded area represent 99% credible bands around the predicted probabilities. The dotted line denotes the threshold of 0.38, used in the classification scheme.

Figure 3: Left Panel: Boxplots showing the probabilities assigned to patients who did achieve a second remission and those who did not for both the Discriminant model and the cox model. Right Panel: Corresponding ROC plot for the Discriminant model (solid red curve) and the cox model (dashed blue curve).

Table 1: Patient Demographics for patients who have and have not experienced a breakthrough seizure and have been observed for long enough to determine their two-year status.

|  |  |  |
| --- | --- | --- |
| **Characteristic** | **Patients who achieve 12-month remission within 2 years of breakthrough seizure** | **Patients who do not achieve 12-month remission within 2 years of breakthrough seizure** |
| **Total**  **(n=185)** | **Total**  **(n=115)** |
| Male | 114 (62%) | 61 (53%) |
| Epilepsy in first degree relative | 26 (14%) | 15 (13%) |
| Neurological insult | 31 (17%) | 19 (17%) |
| **Epilepsy type**  Focal  Generalised  Unclassified | 105 (57%)  51 (28%)  29 (16%) | 80 (70%)  23 (20%)  12 (10%) |
| **EEG results**  Normal  Abnormal  Not done | 58 (32%)  110 (59%)  17 (9%) | 49 (43%)  61 (53%)  5 (4%) |
| **CT/MRI scan results**  Normal  Abnormal  Not done | 91 (49%)  33 (18%)  61 (33%) | 59 (52%)  28 (24%)  28 (24%) |
| **Drugs attempted to achieve 12-month remission**  One  Two or more | 135 (73%)  50 (27%) | 79 (69%)  36 (31%) |
| Number of tonic-clonic seizures ever until first breakthrough seizure, median (IQR) | 2 (1,5) | 2 (0,6) |
| Total number of seizures before diagnosis, median (IQR) | 10 (3,51) | 20 (5,100) |
| Age at first breakthrough seizure, median (IQR) | 24 (16,44) | 35 (20,50) |
| Time to achieve 12-month remission from randomisation (years), median (IQR) | 1 (1,1.52) | 1.24 (1.0,2.0) |
| **Breakthrough seizure treatment decision**  No change to treatment plan  Increase dosage  Decrease dosage (or not specified) | 123 (66%)  59 (32%)  3 (2%) | 60 (52%)  52 (45%)  3 (3%) |

Table 2: Model fixed-effects parameters for the multivariate mixed model.

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
|  | | Longitudinal Variable | | | |
| Seizures since last visit (Yes/No) | | Total Number of Seizures since last visit | |
| Patient Group | Variable | Odds Ratio1 | 95% CI | Parameter Estimate2 | 95% CI |
| Patients who achieve second remission | Time since last follow up (months) | 2.657 | (2.002, 3.495) | 0.024 | (0.012, 0.036) |
| Time since Breakthrough (months) | 0.426 | (0.330, 0.543) | -0.069 | (-0.080, -0.057) |
| Drugs attempted to achieve first remission |  |  | 0.207 | (0.116, 0.302) |
| Age at Breakthrough (Years) |  |  | 0.002 | (-0.001, 0.005) |
| Neurological Insult | 3.739 | (1.147, 11.928) |  |  |
| Patients with no second remission observed | Time since last follow up (months) | 1.487 | (1.291,1.730) | 0.045 | (0.021, 0.070) |
| Time since Breakthrough (months) | 0.819 | (0.769, 0.872) | -0.021 | (-0.039, -0.030) |
| Age at Breakthrough (Years) | 1.030 | (1.010, 1.053) | 0.006 | (-0.004, 0.014) |
| 1. Odds ratios represent the predicted increase/reduction in the odds of experiencing seizures for a given covariate (per one unit increase in continuous covariates, or due to the presence of a binary covariate) 2. Parameter estimates relate to the predicted increase/reduction of the total number of seizures for a given covariate (per one unit increase in continuous covariates, or due to the presence of a binary covariate). 3. Blank entries show that the variable was not included in the submodel for the longitudinal variable described in that column. | | | | | |

Table 3: Prediction Accuracy of the discriminant analysis models and predictions at baseline from a Cox proportional hazards model. The accuracies recorded are the averages across 100 splits of the data into training and test sets.

|  |  |  |
| --- | --- | --- |
|  | Longitudinal Discriminant Analysis | Cox Model |
| Optimal Cutoff | 0.38 | 0.37 |
| Sensitivity | 0.73 | 0.62 |
| Specificity | 0.84 | 0.66 |
| PCC | 0.80 | 0.64 |
| AUC | 0.87 | 0.66 |
| PPV | 0.73 | 0.53 |
| NPV | 0.85 | 0.74 |
| Percentage Unclassified (%) | 17 | 0 |
| Mean Lead Time (days) | 372 | 730 |
| Mean Prediction Time (days) | 303 | 0 |