

An Artificial Intelligence-based Support Tool for Lumbar Spinal Stenosis Diagnosis from Self-Reported History Questionnaire

Frederik Abel^a MD*, Eugene Garcia^{b*}, Vera Andreeva^c MD, Nikolay Nikolaev^{c,d} MD, Serhii Kolisnyk^e MD PhD, Ruslan Sarbaev^b, Ivan Novikov^b BS, Evgeniy Kozinchenko^b, Jack Kim^b PhD, Andrej Rusakov^b MS, Raphael Mourad PhD^{f,b}, Darren R. Lebl MD^a

^aHospital for Special Surgery, 535 East 70th Street, New York, NY 10021, USA

^bRemedy Logic, 1177 Avenue of the Americas, 5th Floor New York, NY, 10036, USA

^cFederal State Budgetary Institution «Federal Center for Traumatology, Orthopedics and Arthroplasty» of the Ministry of Health of the Russian Federation (Cheboksary), Cheboksary, Russia, 428020

^dFederal State Budgetary Educational Institution of Higher Education «Chuvash State University named after I.N. Ulyanov», Cheboksary, Russia, 428015

^eVinnitsa National Medical University, Pyrohova St, 56, Vinnytsia, Vinnytsia Oblast, Ukraine, 21018

^fUniversity of Toulouse, CNRS, UPS, 31062 Toulouse, France

*Co-first authors

Mail addresses:

Frederik Abel, MD: abelf@hss.edu

Eugene Garcia: eugene.garcia@remedylogic.com

Vera Andreeva, MD: Vbarieva@mail.ru

Nikolay Nikolaev, MD: nikolaevns@mail.ru

Serhii Kolisnyk, MD PhD: s.p.kolisnyk@vnmu.edu.ua

Ruslan Sarbaev: ruslan.sarbaev@remedylogic.com

Ivan Novikov: ivan.novikov@remedylogic.com

Evgeniy Kozinchenko: evgeniy.kozinchenko@remedylogic.com

Jack Kim, PhD: jack.kim@remedylogic.com

Andrej Rusakov, MS: a.rusakov@remedylogic.com

Raphael Mourad, PhD: raphael.mourad@univ-tlse3.fr

Darren R. Lebl, MD: research@leblspinemd.com

Corresponding author:

Raphael Mourad, PhD, Université Toulouse III Paul Sabatier, 118 Rte de Narbonne, Toulouse 31062, France,

Email: raphael.mourad@univ-tlse3.fr

Key words:

lumbar spinal stenosis; diagnosis; artificial intelligence; machine learning; self-reported questionnaire.

Running Title:

AI-Support for Lumbar Stenosis Diagnosis

Declaration of Conflicting Interests:

Eugene Garcia, Ruslan Sarbaev, Ivan Novikov, Evgeniy Kozinchenko, Jack Kim, Andrej Rusakov, and Raphael Mourad are employees of Remedy Logic. The rest of the authors declare no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

Source of Funding:

R.M. was supported by Université Paul Sabatier and Remedy Logic. S.K. was supported by Vinnitsa National Medical University and Remedy Logic. J.K. and A.R. were supported by Remedy Logic.

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Abstract

Objectives

Symptomatic lumbar spinal stenosis (LSS) leads to functional impairment and pain. While radiological characterization of the morphological stenosis grade can aid in the diagnosis, it may not always correlate with patient symptoms. Artificial intelligence (AI) may diagnose symptomatic LSS in patients solely based on self-reported history questionnaires.

Methods

We evaluated multiple machine learning (ML) models to determine the likelihood of LSS using a self-reported questionnaire in patients experiencing low back pain and/or numbness in the legs. The questionnaire was built from peer-reviewed literature and a multidisciplinary panel of experts. Random forest, lasso logistic regression, support vector machine, gradient boosting trees, deep neural networks, and automated machine learning models were trained and performance metrics compared.

Results

Data from 4,827 patients (4,690 patients without LSS: mean age 62.44, range 27 – 84 years, 62.8% females, and 137 patients with LSS: mean age 50.59, range 30 – 71 years, 59.9% females) were retrospectively collected. Among the evaluated models, the random forest model demonstrated the highest predictive accuracy with an area under the receiver operating characteristic curve (AUROC) between model prediction and LSS diagnosis of 0.96, a sensitivity of 0.94, a specificity of 0.88, a balanced accuracy of 0.91 and a Cohen’s kappa of 0.85.

Conclusions

Our results indicate that ML can automate the diagnosis of LSS based on self-reported questionnaires with high accuracy. Implementation of standardized and intelligence-automated workflow may serve as a supportive diagnostic tool to streamline patient management and potentially lower healthcare costs.

28

29 **Introduction**

30 Degenerative lumbar spinal stenosis (LSS) is a significant source of disability in older adults,
31 which affects an estimated 103 million persons annually worldwide¹. The clinical syndrome of LSS is
32 characterized by chronic lower back and extremity pain, accompanied by loss of mobility and function,
33 which can steadily reduce patients' quality of life². First-line treatments include modification of activity,
34 analgesia, and physical therapy. In cases where conservative treatments fail, decompressive spinal surgery
35 is often considered as an option to relieve symptoms. As such, LSS is associated with a significant
36 socioeconomic burden and high healthcare costs^{3,4}. Hence, strategies for simplified, fast, and automated
37 diagnosis based on clinical symptoms may have an important societal impact.

38 Substantial improvements of standardized diagnostic accuracy are streamlined by artificial
39 intelligence (AI), in particular machine learning (ML) models to develop data-driven algorithms and
40 intelligent automation (IA) to enhance human intelligence and therapeutic-decision making. Most
41 commonly, ML models have successfully been trained to automatically detect and classify LSS based on
42 MRI studies of the lumbar spine, achieving high accuracy levels comparable to those of subspecialist
43 radiologists^{5,6}. Other approaches applied ML to determine surgical candidacy for spinal surgery based on
44 lumbar spine MRI's⁷ or by using hybrid AI models, that combine features from both imaging and clinical
45 information⁸. More recently, ML methods have been utilized to determine prior authorization approval for
46 LSS surgery based on medical vignettes, which consisted of both clinical data and MRI findings⁹.
47 Although diagnostic imaging is a mainstay for the evaluation of LSS, radiographically affected patients
48 may be clinically asymptomatic. Therefore, clinical symptoms also play a significant role in therapeutic
49 decision-making. Patients with symptomatic LSS can be assessed using self-reported questionnaires¹⁰.
50 However, ML has not yet been tested to identify such patients based solely on patient questionnaires
51 without any imaging data, which could potentially facilitate intelligence-automated therapeutic decision-
52 making.

53 Here, we propose a novel AI approach to diagnose LSS from self-reported history questionnaires
54 that assess clinical history, pain character, and mobility of patients. Different ML models were trained on

55 retrospectively collected data from patients with diagnosed LSS (LSS+) and without LSS (LSS-) and their
56 performance compared, including random forest, lasso logistic regression, support vector machine
57 (SVM), gradient boosting trees, deep neural networks (DNN), and automated machine learning (H2O
58 autoML). Finally, key contributor variables for predicting patients with LSS+ were identified.

59

60 **Materials and Methods**

61

62 *Subjects and Data collection*

63 This multicenter study was performed in two hospitals (BLINDED FOR REVIEW and
64 BLINDED FOR REVIEW), and one outpatient clinic (BLINDED FOR REVIEW). Data of patients were
65 retrospectively collected from self-reported questionnaires from August 2021 to September 2022.
66 Additionally, health records were assessed, including past clinical history, treatments, and results of
67 examinations, for all patients. The selection was limited to patients who presented with primary
68 symptoms of low back pain and/or numbness in the back and legs, and experienced difficulties in
69 performing daily activities. The age criteria for inclusion were set at 20 years or older. The questionnaires
70 were filled out by our clinical administrators based on the answers provided by each interviewed patient.
71 LSS diagnosis (presence/absence) was confirmed by an orthopedic surgeon based on clinical history of
72 each patient and reports from lumbar spine MRI studies, which served as the ground truth.

73 This retrospective study received institutional review board approval and written informed
74 consent was obtained from all subjects.

75

76 *Questionnaire*

77 A literature review was performed to identify peer-reviewed medical literature that assess
78 diagnosis and outcome measures of LSS (Supplementary material 1). Items from these articles, as well as
79 relevant items extracted from The Short Form (36) Health Survey (SF-36)¹¹, EQ-5D¹², and Oswestry
80 Disability Index (ODI)¹³, were collected. In total, 205 questions were accumulated, which were then
81 compiled with the input of an expert panel comprising a multidisciplinary team of doctors in the fields of

82 spinal surgery, rehabilitation medicine, interventional and diagnostic radiology. The final self-reported
83 questionnaire included 26 questions (qualitative and continuous outcome variables) including pain, pain
84 severity and type, activities prevented by pain (e.g. pain prevents sleeping rate), but also motor
85 impairment, the use of moving device, history of spinal cord or cauda equina injury, general health, and
86 mental health (e.g. anxiety or depression level) (**Table 1**).

87

88 *Data Imputation and Machine Learning*

89 The dataset comprised 4,690 patients without LSS (LSS-; mean age 62.44; range 27 – 84 years;
90 62.8% females) and 137 patients with LSS (LSS+; mean age 50.59; range 30 – 71 years: 59.9% females),
91 summing up to 4,827 patients in total. The overall percentage of missing data was 29%. To cope with
92 missing data, values were imputed applying the median, where necessary. Subsequently, the data were
93 used to train the machine learning models. Patient data were randomly split into 80% for fine-tuning and
94 training of the machine learning models (3,758 LSS- and 104 LSS+) and 20% for testing predictions (932
95 LSS- and 33 LSS+). To ensure a balanced training dataset for effective model training, patients without
96 LSS were down-sampled to match the number of patients with LSS, resulting in a total of 208 patients for
97 training (104 LSS- and 104 LSS+). Different machine learning models were trained and compared,
98 including random forest, lasso logistic regression, support vector machine (SVM), gradient boosting trees
99 (XGBOOST), deep neural network (DNN), and automated machine learning (H2O autoML). Variable
100 importance was computed using the mean decrease in accuracy in the out-of-bag sample during training.
101 Hyper-parameters were obtained by fine-tuning with 5-fold cross-validation.

102

103 *Data and Statistical Analysis*

104 All data analyses, including univariate and bivariate analyses, prediction performance metrics,
105 and plots were done using R (Version 4.2.1, the R Foundation). The following R packages were used for
106 computations and fine-tuning: ranger for random forest and variable importance ([https://cran.r-](https://cran.r-project.org/web/packages/ranger)
107 [project.org/web/packages/ranger](https://cran.r-project.org/web/packages/ranger)), tuneRanger for hyper-parameter fine-tuning ([https://cran.r-](https://cran.r-project.org/web/packages/tuneRanger/)
108 [project.org/web/packages/tuneRanger/](https://cran.r-project.org/web/packages/tuneRanger/)), glmnet for lasso logistic regression ([https://cran.r-](https://cran.r-project.org/web/packages/glmnet/)

109 project.org/web/packages/glmnet), e1071 for SVM (<https://cran.r-project.org/web/packages/e1071>),
110 XGBOOST for extreme gradient boosting (<https://cran.r-project.org/web/packages/xgboost>), nnet for
111 deep neural network (<https://cran.r-project.org/web/packages/nnet>), and caret for SVM, XGBOOST, and
112 DNN fine-tuning (<https://topepo.github.io/caret/>). For quantitative variables, differences of means
113 between the LSS+ patients and LSS- patients were tested using the Student's test. For qualitative
114 variables, differences of proportions between the LSS+ patients and LSS- patients were assessed using
115 the Fisher's exact test. To account for multiple tests, the Bonferroni p -value threshold was used and
116 computed as $0.05/34=0.0014$. Performance metrics, including area under the receiver operating
117 characteristic curve (AUROC), area under the precision-recall curve (AUPRC), sensitivity, specificity,
118 Cohen's kappa, accuracy, and the F1-score, were calculated for all models.

119

120 **Results**

121

122 *Univariate analysis of LSS predictors*

123 **Table 2** displays the differences in assessed outcome variables between patients with LSS
124 (LSS+) and patients without LSS (LSS-). Most predictors exhibited significant p -values, with $p<0.0014$
125 (Bonferroni threshold). It included predictors describing general health (mean difference of -18 points out
126 of 100, 95% confidence interval (CI) [-20.71; -15.45], $p<0.0001$), mental health (mean difference of -17
127 points out of 100, 95%CI [-19.98; -14.34], $p<0.0001$), pain severity (mean difference of +25.37 points out
128 of 100, 95%CI [22.64; 28.11], $p<0.0001$), or pain preventing activities, e.g. pain preventing standing
129 (mean difference of +35.7 points out of 100, 95%CI [31.41; 39.99], $p<0.0001$). Additionally, reduced or
130 damaged motor skills, problems with performing daily activities (i.e. problems performing washing or
131 dressing), or use of moving devices were significantly different between LSS+ and LSS- patients. Thus,
132 univariate analysis demonstrated strong associations between LSS and most self-reported history
133 predictors, suggesting their potential use in building a ML model for accurate prediction of LSS.

134

135 *Prediction of LSS based on Machine Learning*

136 The accuracy of our ML approach in predicting LSS was assessed and compared with the ground
137 truth. For this purpose, the patient history data were randomly split into 80% of patients for training a
138 series of ML models, while the remaining 20% of patients were reserved to estimate the prediction
139 accuracy of the different models. The performance of the random forest, lasso regression, SVM,
140 XGBOOST, DNN, and automated machine learning models is summarized in **Table 3**.

141 Most ML models showed excellent prediction performances when classifying LSS+ versus LSS-
142 patients. Among the models, the random forest exhibited the highest prediction performance, achieving
143 an AUROC of 0.96 (95% CI [0.949; 0.980]), sensitivity of 0.94, specificity of 0.88, and Cohen's kappa
144 value of 0.85 (**Figure 1A**). The second-best performing model was XGBOOST, which demonstrated an
145 AUROC of 0.96, a sensitivity of 0.97, a specificity of 0.86, and a Cohen's kappa value of 0.88 (**Figure**
146 **1D**).

147 Since the data were highly imbalanced (4,690 LSS+ patients and 137 LSS- patients), we also
148 computed the balanced accuracy, a more suitable metric for imbalanced data. The random forest also
149 showed a high balanced accuracy of 0.91, while the XGBOOST model achieved the highest value of
150 0.92. Our random forest was chosen as the optimal trade-off between a high AUROC and balanced
151 accuracy for predicting LSS from self-reported history data, demonstrating excellent predictive
152 performance metrics.

153

154 *Importance of predictors*

155 Next, we conducted an assessment to identify the key predictors of LSS. For this purpose, we
156 computed variable importance using the random forest model to identify the most significant predictors
157 (**Figure 2**). Among the list of predictors, problems with performing daily activities, including washing or
158 dressing, were found to be the most significant predictors for LSS+ patients. Additionally, pain severity,
159 and pain or emotional distress that restricts social interactions and/or activities were significant
160 contributors to patients with LSS.

161

162 **Discussion**

163

164 Applications of artificial intelligence (AI), including machine learning (ML) models and
165 intelligent automation (IA), are increasing rapidly in the medical domain and have demonstrated
166 remarkable success in various clinical settings and research areas. AI systems strive to improve diagnostic
167 processes, prognostication, and outcomes in a transparent and observer-independent manner, thereby
168 enhancing therapeutic decision-making. The field of spine surgery, in particular, benefits from AI
169 applications and IA tools, as diagnostics and therapeutic decision-making often require clinical expertise
170 and rely on interpretation of ramified factors such as medical history, imaging, or perioperative data¹⁴.

171 In this article, we tested a series of ML models to predict symptomatic LSS based on a simple
172 self-reported history questionnaire of patients with low back pain and/or numbness in the back and legs,
173 and having problems performing daily activities. Data from 4,827 patients (4,690 LSS- and 137 LSS+)
174 were collected and key factors identified that revealed strong association with LSS+ patients.
175 Subsequently, different ML models were trained and evaluated on a balanced subset of the self-reported
176 predictors, including random forest, lasso logistic regression, support vector machine (SVM), gradient
177 boosting trees, deep neural networks (DNN), and H2O automated machine learning. Of these, the random
178 forest model demonstrated the highest diagnostic accuracy with a prediction error as measured by the
179 AUROC of 0.96, a sensitivity of 0.94, a specificity of 0.88, a balanced accuracy of 0.91, and a Cohen's
180 kappa of 0.85. The computation of variable importance revealed that problems with performing daily
181 activities, pain severity and emotional distress that restrict social interaction or activity rate as were the
182 most significant contributors to patients with LSS+. As such, self-reported questionnaires may be feasible
183 to predict symptomatic LSS patients in an IA-based manner, ultimately enhancing human therapeutic
184 decision-making.

185 Previous studies have utilized AI applications in spine surgery, with a focus on radiological
186 features extracted from MRI data and employing various ML models to detect and diagnose LSS^{5,6,15}.
187 While radiological characterization of morphological stenosis grade contributes to the diagnosis of LSS, it
188 may not always correlate with pain intensity and functional disability experienced by affected
189 patients^{16,17}. Consequently, incorporating AI-guided diagnosis of LSS based on other aspects or in

190 combination with radiological features may be a more comprehensive approach to reflect the clinical
191 syndrome of LSS. However, such approaches have been investigated in only a limited number of studies.
192 Ren *et al.* investigated natural language processing-based ML models based on positive symptoms
193 extracted from electronic health records¹⁸. Contrasting our study, different models were tested to
194 discriminate patients with LSS from patients with lumbar disc herniation, and found that a Long Short-
195 Term Memory DNN achieved the highest capacity with an AUROC of 0.85 for this task. Another
196 approach used ML algorithms on data of patients that performed five-repetition sit-to-stand tests¹⁹. The
197 algorithm, a fuzzy rule-based system, achieved a classification accuracy of 96.2% for patients with disc
198 herniation, LSS, and chronic lower back pain. Five-repetition sit-to-stand tests are designed to assess
199 functional impairment, which has been identified as the most significant contributor variable to LSS+
200 patients in our dataset, followed by pain and emotional distress that restrict social interaction or activity
201 rate.

202 Accordingly, these predictors have been determined as critical factors for treatment and decision-
203 making in LSS+ patients by previous studies. The presence of disability, along with pain and radiological
204 stenosis grade, has been associated with the likelihood of requiring surgical therapy²⁰. Similarly,
205 functional disability and pain severity have been correlated with impairment of health-related quality of
206 life in patients with LSS or lumbar disc herniation²¹. Pain severity has been as identified as the fourth
207 most important contributor to LSS patients in our dataset, suggesting that actual restriction of daily
208 activities may be more significant for the diagnosis of LSS patients. However, it should be considered
209 that pain and functional impairment are closely related, and the inability to perform daily activities likely
210 is a secondary effect of pain. Another significant predictive variable for the diagnosis of LSS was the
211 patient's mental status, evaluated through our questionnaire based on the patient's subjective grading of
212 their mental health, encompassing a broader field of psychological factors including depression or
213 anxiety. Prior studies have indicated that these factors, particularly preoperative depression, are associated
214 with increased severity of postoperative LSS-related symptoms and poorer long-term outcomes following
215 decompression^{22,23}. Therefore, although it can be challenging to assess in clinical practice, considering the

216 patient's overall mental health could significantly contribute to a more comprehensive management
217 approach and improve the patient's prognosis.

218 Our study contains several limitations. First, the dataset was comprised 4690 LSS- patients and
219 only 137 LSS+ patients, which required down-sampling to obtain a more balanced sample size. Second,
220 the ground truth of LSS+ diagnosis in this cohort was based on clinical history of each patient and reports
221 from MRI examinations and did not include independent classification by other experts. Third, our ML
222 models solely focused on diagnosis of LSS on basis of self-reported symptoms in questionnaires.
223 Symptoms of LSS may partially overlap those of other concomitant degenerative spinal disorders (e.g.
224 degenerative disc disease, spondylolisthesis) in these patients and differentiation of these entities was not
225 addressed by our models. Additionally, other factors relevant for management of LSS were not
226 investigated, including surgical decision-making or impairment of patients' quality of life. Fourth, our
227 algorithms were not validated in an external patient cohort within this study, potentially limiting their
228 generalizability. Finally, our proposed approach is simplified and did not integrate other features, such as
229 radiological stenosis grade, that may have further increased diagnostic accuracy in detecting LSS patients.

230

231 **Conclusions**

232

233 In summary, our results demonstrate that AI can be applied to diagnose symptomatic LSS in
234 patients based on simplified, self-reported history questionnaires with high accuracy even in the absence
235 of any imaging input into the model. Functional impairment and pain/emotional distress that restrict
236 social interaction or activity rate are key contributors to patients with LSS. Implementation of
237 standardized and automated AI-guided workflow may act as an intelligent automation tool to identify
238 patients with LSS using simple self-reported history questionnaires and may more efficiently and cost-
239 effectively help determine which patients required advanced imaging studies such as MRI and
240 consideration for surgery.

241

242

243 **Acknowledgements**

244 N/A

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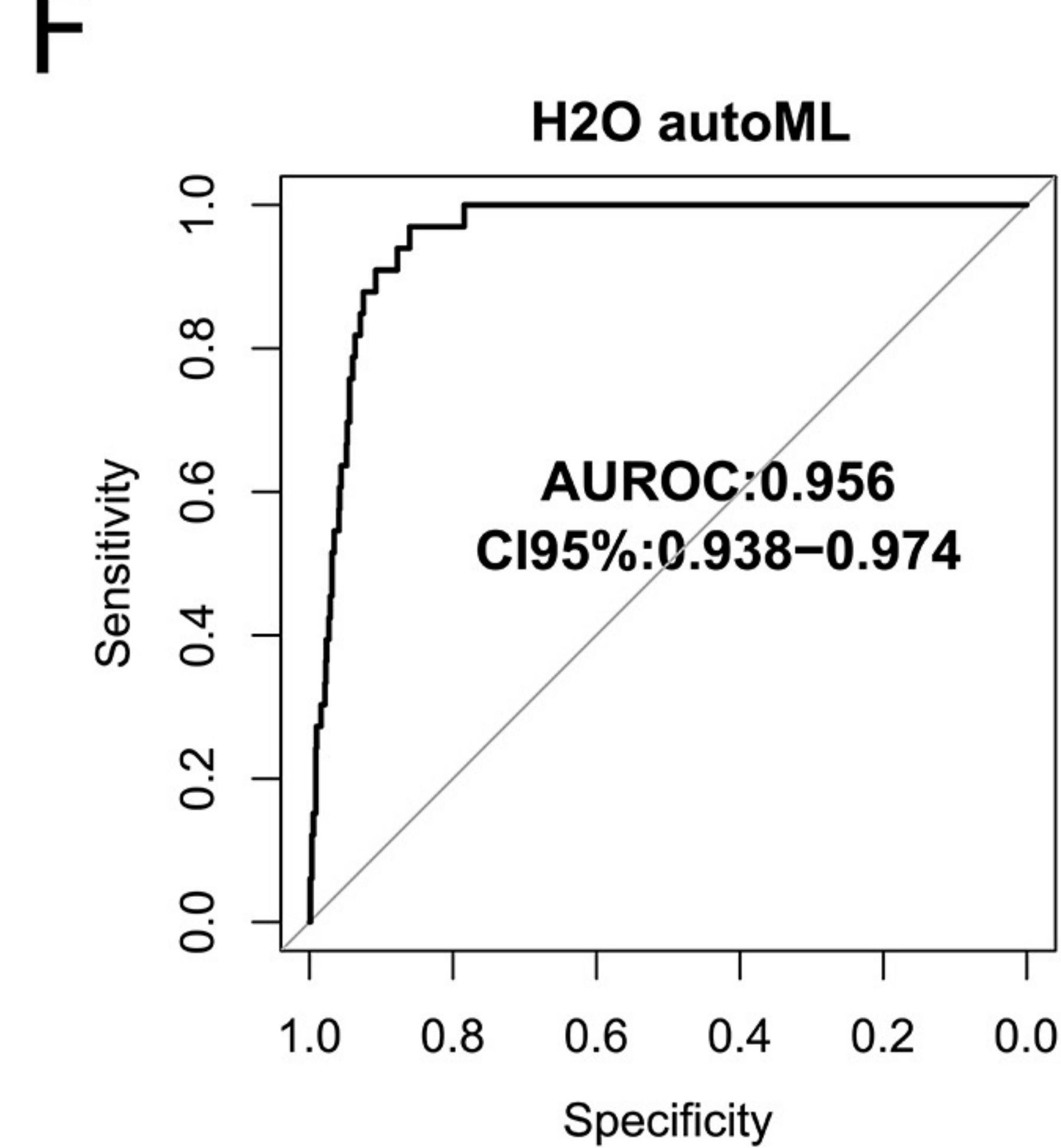
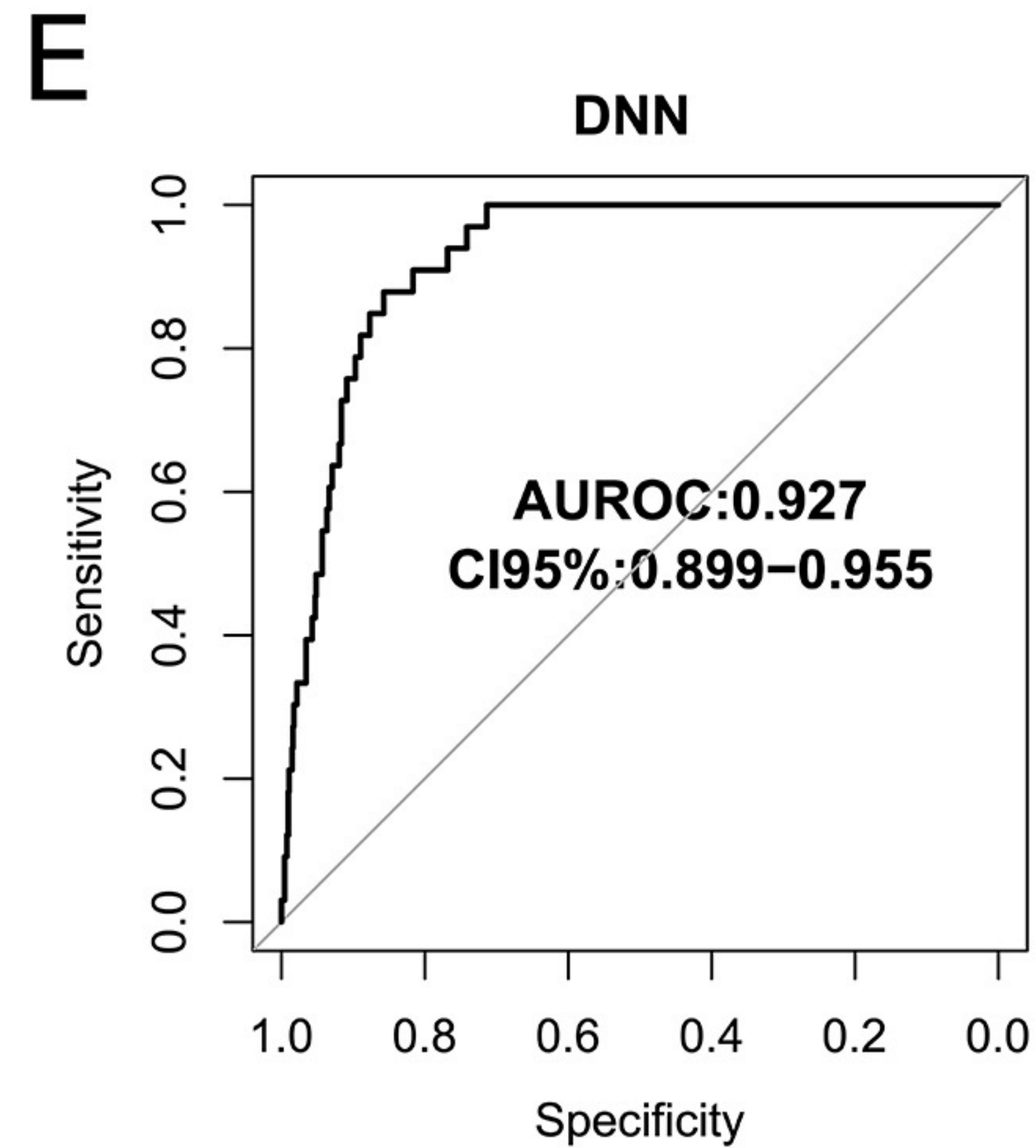
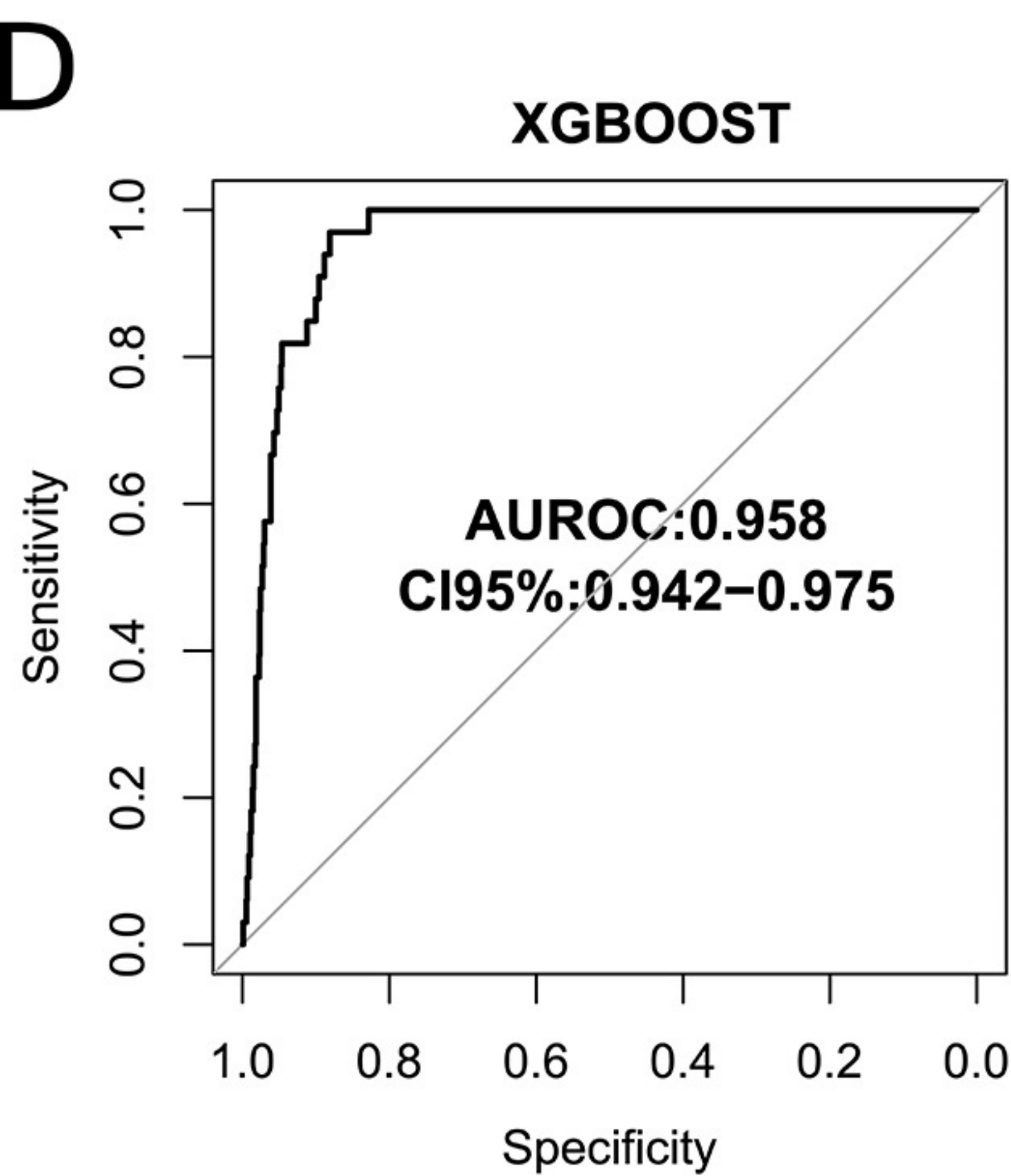
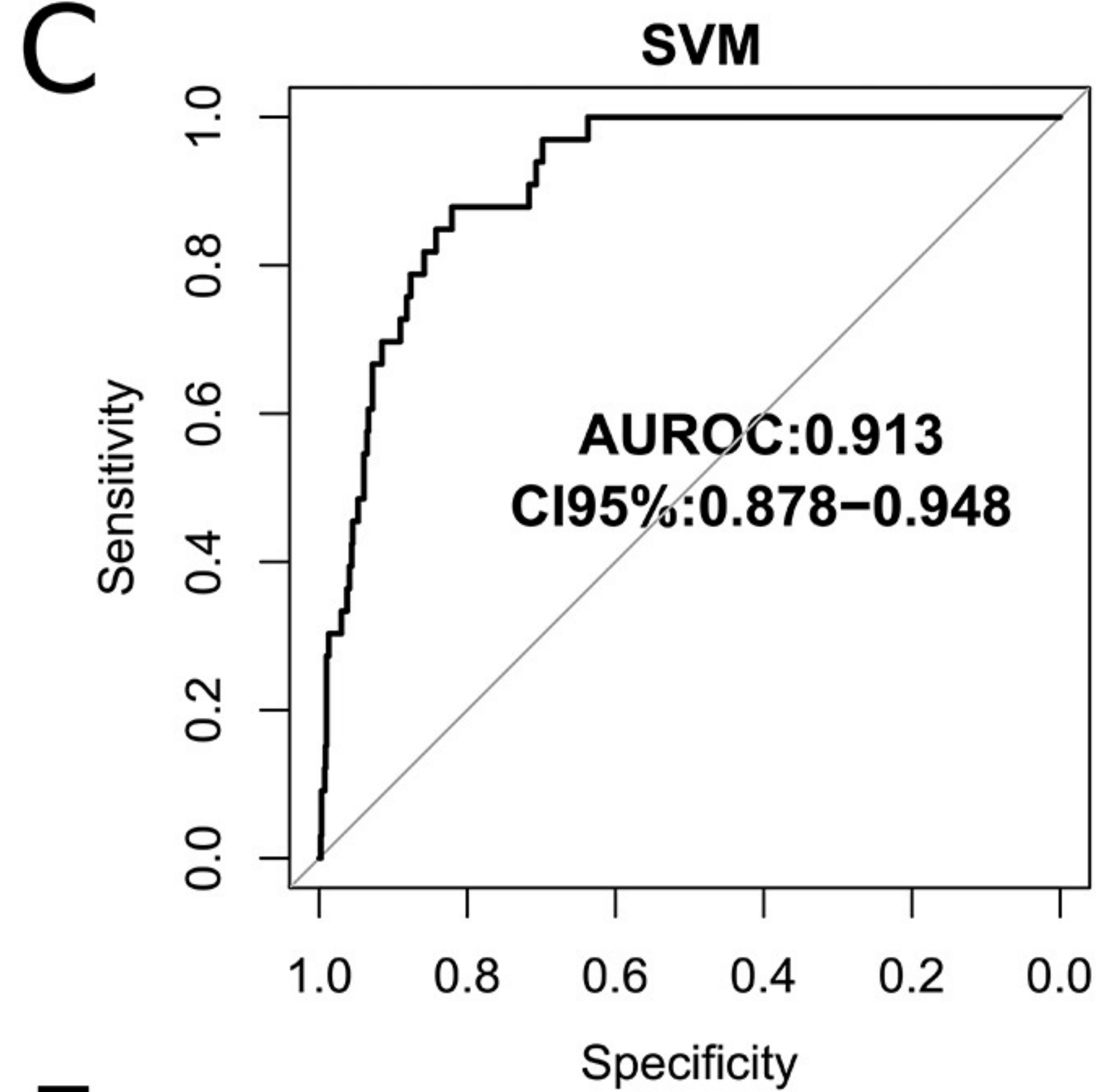
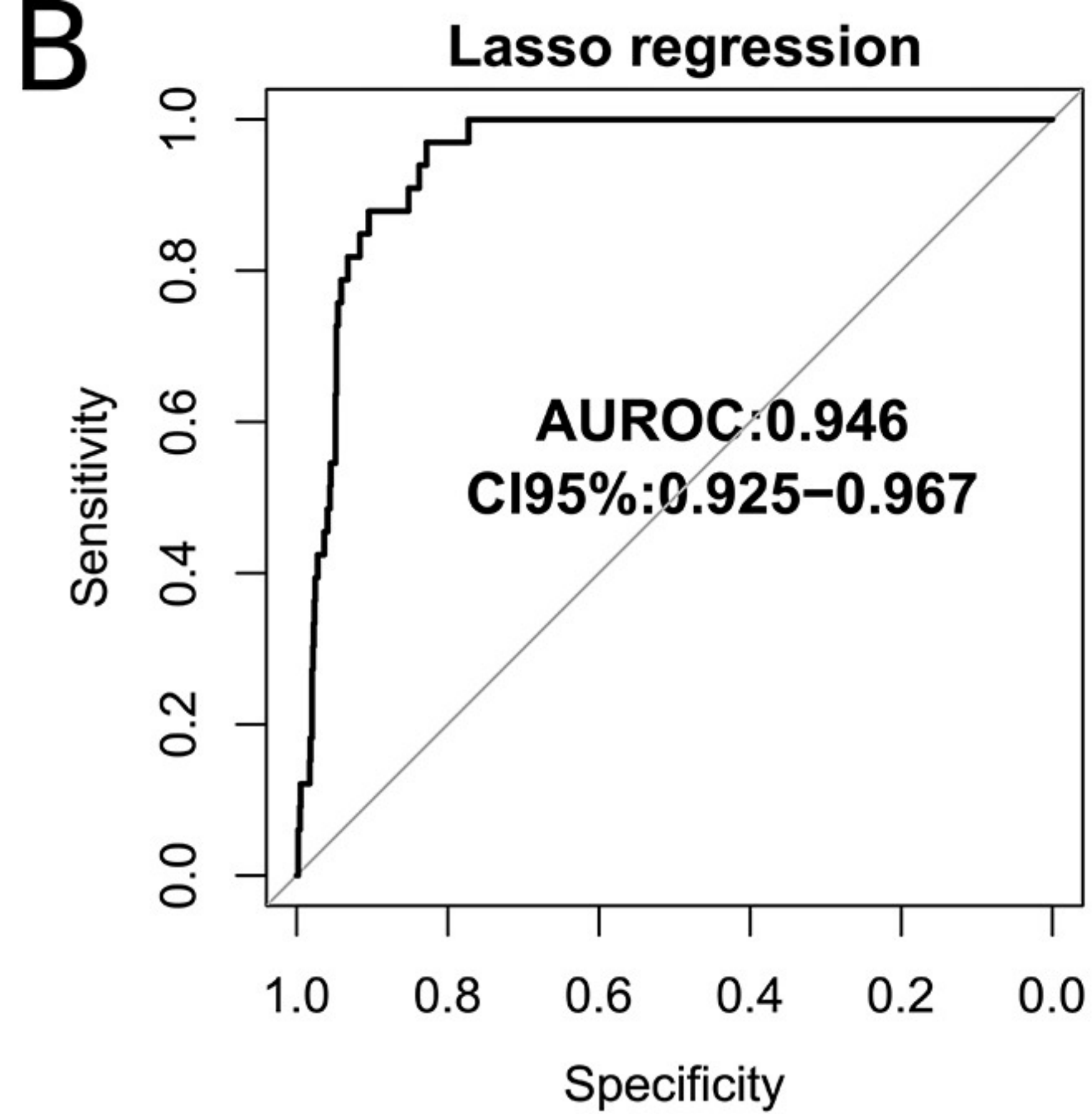
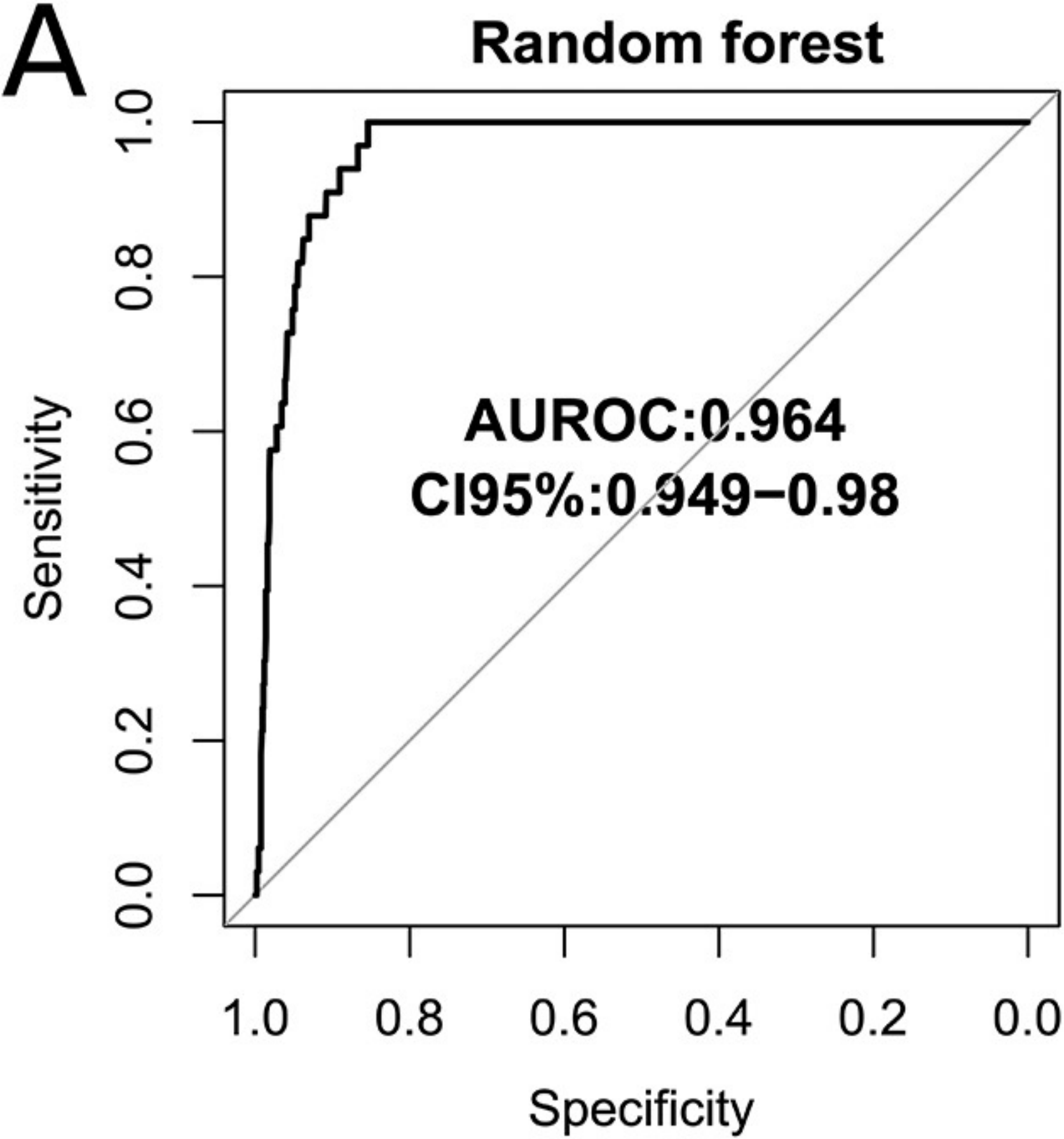
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309 **Figure Legends**
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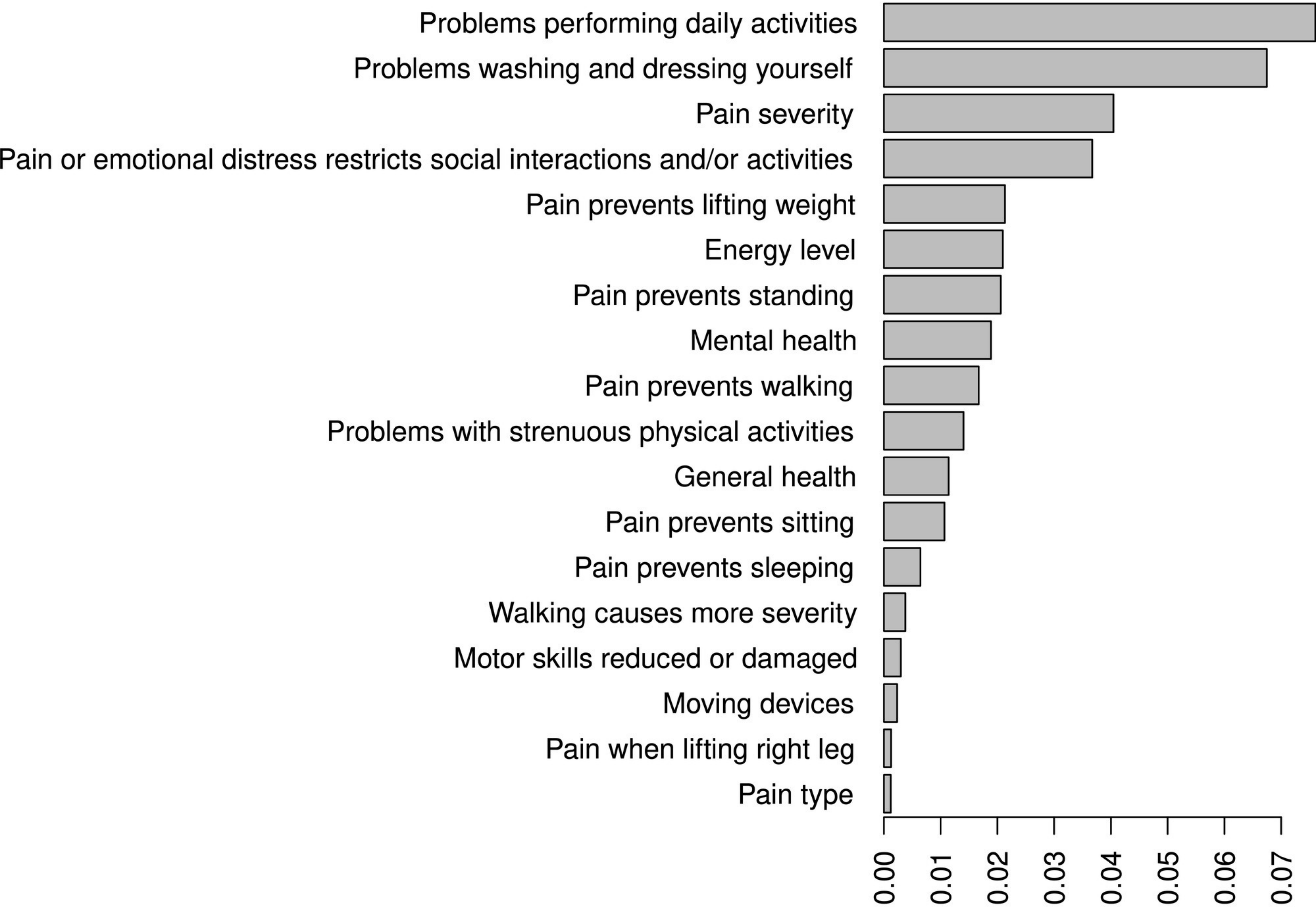
311 **Figure 1:** Prediction performance of lumbar spinal stenosis (LSS) based on self-reported history
312 questionnaires for several machine learning models. Receiver operating characteristic curve (ROC) of the
313 random forest (A), lasso logistic regression (B), support vector machine (SVM, C), XGBOOST (D), deep
314 neural network (DNN, E), and automated machine learning (H2O autoML, F).

315
316 **Figure 2:** Top key variables to predict lumbar spinal stenosis probability. Variable importance was
317 calculated using random forests with permutations.

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Variable Importance



Tables

Table 1: Self-reported questionnaire with 26 questions to assess manifestation of symptomatic LSS.

| Question | Value |
|---|---------|
| 1. Lumbar spinal stenosis | Yes/No |
| 2. General health (0 – Worst; 100 – Best general health imaginable) | 0 – 100 |
| 3. Mental health (0 – Worst; 100 – Best mental health imaginable) | 0 – 100 |
| 4. Congenital diseases or conditions | Yes/No |
| 5. Spinal cord or cauda equina injury | Yes/No |
| 6. Pain severity (0 – Zero pain; 100 – Extreme pain) | 0 – 100 |
| 7. Pain type | |
| Extreme cold | Yes/No |
| Extreme hot or burning sensation | Yes/No |
| Itching | Yes/No |
| Mechanical | Yes/No |
| Sharp stabbing | Yes/No |
| Throbbing and/or pulsating | Yes/No |
| 8. Bending forward increases pain severity | Yes/No |
| 9. Standing up increases pain severity | Yes/No |
| 10. Walking increases pain severity | Yes/No |
| 11. Pain increases while sleeping | Yes/No |
| 12. Pain prevents lifting weights (0 – Zero pain; 100 – Extreme pain) | 0 – 100 |
| 13. Pain prevents sitting (0 – Zero pain; 100 – Extreme pain) | 0 – 100 |
| 14. Pain prevents sleeping (0 – Zero pain; 100 – Extreme pain) | 0 – 100 |
| 15. Pain prevents standing (0 – Zero pain; 100 – Extreme pain) | 0 – 100 |
| 16. Pain prevents walking (0 – Zero pain; 100 – Extreme pain) | 0 – 100 |
| 17. Daily activities limited by physical pain | Yes/No |
| 18. Pain when lifting right leg: | |
| - No | Yes/No |
| - Back pain only | Yes/No |
| - Pain radiating down the leg | Yes/No |
| 19. Pain when lifting left leg: | |
| - No | Yes/No |
| - Back pain only | Yes/No |
| - Pain radiating down the leg | Yes/No |
| 20. Motor skills reduced or damaged | Yes/No |
| 21. Moving devices | Yes/No |
| 22. Problems performing daily activities (0 – Zero problems; 100 – Extreme problems) | 0 – 100 |
| 23. Problems washing and dressing yourself (0 – Zero problems; 100 – Extreme problems) | 0 – 100 |
| 24. Problems with strenuous physical activities (0 – Zero problems; 100 – Extreme problems) | 0 – 100 |
| 25. Energy level (0 – No energy at all; 100 – Highest energy level imaginable) | 0 – 100 |
| 26. Pain or emotional distress (0 – Zero; 100 – Extreme pain/emotional distress) | 0 – 100 |

Table 2: Univariate analyses of predictors associated with the diagnosis of LSS.

| Variable | LSS- (n=4644) | LSS+ (n=137) | OR | Mean difference | 95%CI | p-value |
|---|------------------|-----------------|-------|-----------------|------------------|---------|
| General health | 57.6 | 39.51 | | -18 | [-20.71; -15.45] | <0.0001 |
| Mental health | 66.83 | 49.67 | | -17 | [-19.98; -14.34] | <0.0001 |
| Congenital diseases/conditions | 11.68% | 36.50% | 4.34 | | [2.97; 6.30] | <0.0001 |
| Spinal cord or cauda equina injury | 14.71% | 50.36% | 5.88 | | [4.10; 8.43] | <0.0001 |
| Pain severity | 46.09 | 71.46 | | 25.37 | [22.64; 28.11] | <0.0001 |
| Pain type | | | | | | |
| - Extreme cold | 0.23% | 0.73% | 3.08 | | [0.07; 21.51] | 0.2959 |
| - Extreme hot or burning sensation | 7.06% | 15.33% | 2.37 | | [1.39; 3.87] | 0.0012 |
| - Itching | 0.58% | 0.73% | 1.25 | | [0.03; 7.72] | 0.5596 |
| - Mechanical | 18.42% | 14.60% | 0.74 | | [0.43; 1.21] | 0.2604 |
| - Sharp stabbing | 18.72% | 27.74% | 1.66 | | [1.09; 2.47] | 0.0141 |
| - Throbbing and/or pulsating | 6.20% | 5.11% | 0.8 | | [0.31; 1.72] | 0.7198 |
| Bending forward increases pain severity | 18.61% | 32.12% | 2.07 | | [1.40; 3.02] | <0.0001 |
| Standing up increases pain severity | 10.13% | 27.74% | 3.4 | | [2.25; 5.06] | <0.0001 |
| Walking increases pain severity | 14.05% | 48.18% | 5.68 | | [3.96; 8.15] | <0.0001 |
| Pain increases while sleeping | 22.37% | 48.18% | 3.34 | | [2.30; 4.84] | <0.0001 |
| Pain prevents lifting weights | 42.36 | 75.41 | | 33.05 | [30.08; 36.03] | <0.0001 |
| Pain prevents sitting | 32.53 | 57.15 | | 24.62 | [20.76; 28.48] | <0.0001 |
| Pain prevents sleeping | 26.68 | 51.69 | | 25.01 | [20.22; 29.81] | <0.0001 |
| Pain prevents standing | 33.92 | 69.62 | | 35.7 | [31.41; 39.99] | <0.0001 |
| Pain prevents walking | 35.00 | 69.64 | | 34.64 | [30.39; 38.89] | <0.0001 |
| Daily activities limited by physical pain | 82.20% | 98.54% | 14.62 | | [3.95; 122.34] | <0.0001 |
| Pain when lifting right leg: | | | | | | |
| - No | 69.46% | 45.26% | 0.36 | | [0.25; 0.52] | <0.0001 |
| - Back pain only | 10.21% | 18.25% | 1.96 | | [1.20; 3.08] | 0.0044 |
| - Pain radiating down the leg | 4.71% | 15.33% | 3.66 | | [2.14; 5.99] | <0.0001 |
| Pain when lifting left leg: | | | | | | |
| - No | 68.14% | 43.07% | 0.35 | | [0.25; 0.50] | <0.0001 |
| - Back pain only | 10.77% | 19.71% | 2.03 | | [1.27; 3.16] | 0.0021 |
| - Pain radiating down the leg | 4.78% | 15.33% | 3.61 | | [2.11; 5.91] | <0.0001 |
| Motor skills reduced or damaged | 57.38% | 93.43% | 10.56 | | [5.37; 23.68] | <0.0001 |
| Moving devices | 5.48% | 34.31% | 9 | | [6.04; 13.26] | <0.0001 |
| Problems performing daily activities | 25.94 | 59.12 | | 33.18 | [30.20; 36.17] | <0.0001 |
| Problems washing and dressing yourself | 20.77 | 42.54 | | 21.77 | [19.23; 24.32] | <0.0001 |
| Problems with strenuous physical activities | 34.48 | 61.54 | | 27.06 | [23.13; 30.00] | <0.0001 |
| Energy level | 61.49 | 35.56 | | -25.93 | [-29.06; -22.81] | <0.0001 |
| Pain or emotional distress | 24.12 | 59.80 | | 35.68 | [31.67; 39.68] | <0.0001 |

95%CI = 95% confidence interval; LSS = Lumbar spinal stenosis; OR = Odds ratio.

Table 3: Table of predictive performance metrics for the different models.

| Model | Sensitivity | Specificity | Kappa | AUROC | Accuracy | F1 | Balanced Accuracy | AUPRC |
|------------------|-------------|-------------|--------|--------|----------|--------|-------------------|--------|
| RF | 0.9394 | 0.8777 | 0.8485 | 0.9645 | 0.8798 | 0.9338 | 0.9085 | 0.8943 |
| Lasso regression | 0.8788 | 0.8809 | 0.8485 | 0.9461 | 0.8808 | 0.9345 | 0.8798 | 0.9008 |
| SVM | 0.8485 | 0.8273 | 0.7576 | 0.9129 | 0.828 | 0.9028 | 0.8379 | 0.9062 |
| XGBOOST | 0.9697 | 0.8627 | 0.8788 | 0.9582 | 0.8663 | 0.9257 | 0.9162 | 0.8971 |
| DNN | 0.8788 | 0.838 | 0.7576 | 0.9268 | 0.8394 | 0.9097 | 0.8584 | 0.9035 |
| H2O autoML | 0.9394 | 0.8691 | 0.8485 | 0.9556 | 0.8715 | 0.9289 | 0.9042 | 0.896 |

AUPR = Area under precision-recall curve; AUROC = Area under the receiver operating characteristic; DNN = Deep neural network; RF = Random forest; SVM = Support vector machine.