

04/19/2020
 bioRxiv

Simple Behavioral Analysis (SimBA) – an open source toolkit for computer classification of complex social behaviors in experimental animals.

Simon RO Nilsson^{†1,3}, Nastacia L. Goodwin^{†1,2}, Jia Jie Choong^{†1}, Sophia Hwang¹, Hayden R Wright⁴, Zane C Norville⁶, Xiaoyu Tong⁷, Dayu Lin⁷, Brandon S. Bentzley⁶, Neir Eshel⁶, Ryan J McLaughlin^{4,5}, Sam A. Golden^{1,2,3}

1 University of Washington, Department of Biological Structure, Seattle, WA, USA

2 University of Washington, Graduate Program in Neuroscience, Seattle, WA, USA

3 University of Washington, Center of Excellence in Neurobiology of Addiction, Pain, and Emotion (NAPE), Seattle, Washington, USA

4 Washington State University, Department of Integrative Physiology and Neuroscience, Pullman, WA, USA

5 Washington State University, Departments of Psychology, Pullman, WA, USA

6 Stanford University, Department of Psychiatry and Behavioral Sciences, Stanford, CA, USA

7 New York University, Neuroscience Institute, New York, NY

[†]Denotes equal contribution

Running title: Automated classification of rodent social behavior

Text information

Abstract: 311 words

Introduction: 744 words

Methods: 2469 words

Results: 1148 words

Discussion: 1302 words

Total text: 5974 words

Figures: 11

Tables: 4

References: 106

* Corresponding Author: Sam A. Golden (sagolden@uw.edu) and Simon RO Nilsson (snilsson@uw.edu)

Table of contents:

1. Abstract
2. Graphical abstract
3. Introduction
4. Methods
5. Results
6. Discussion
7. Conclusion

Author contributions: SRON and SAG conceived of the software and all authors contributed intellectually to its development. SRON, JJC, and SH wrote the SimBA code. XT wrote the SEQ to MP4 conversion code. SRON, NLG, HRW and ZCN generated the classifiers. JJC, SRON and SAG wrote the online documentation. SRON and SAG created the manuscript figures. NE, BSB and ZCN aligned photometry and social classifications. SRON, NLG and SAG wrote the manuscript with input from all other authors.

Funding and disclosure:

The authors declare that they do not have any conflicts of interest (financial or otherwise) related to the text of the paper. The research was supported by NIDA 4R00DA045662-02 (SAG), NIDA P30 DA048736 (SRON and SAG), NARSAD Young Investigator Award 27082 (SAG), and NIDA T32 5T32NS099578-04 (NLG). We thank Briana Smith, Liana Bloom, Annette Mercedes, Roël Vrooman and Kayla Pitts for their thoughtful assistance annotating behavioral frames and discussing the manuscript and online documentation.

Abstract

Aberrant social behavior is a core feature of many neuropsychiatric disorders, yet the study of complex social behavior in freely moving rodents is relatively infrequently incorporated into preclinical models. This likely contributes to limited translational impact. A major bottleneck for the adoption of socially complex, ethology-rich, preclinical procedures are the technical limitations for consistently annotating detailed behavioral repertoires of rodent social behavior. Manual annotation is subjective, prone to observer drift, and extremely time-intensive. Commercial approaches are expensive and inferior to manual annotation. Open-source alternatives often require significant investments in specialized hardware and significant computational and programming knowledge. By combining recent computational advances in convolutional neural networks and pose-estimation with further machine learning analysis, complex rodent social behavior is primed for inclusion under the umbrella of computational neuroethology.

Here we present an open-source package with graphical interface and workflow (Simple Behavioral Analysis, SimBA) that uses pose-estimation to create supervised machine learning predictive classifiers of rodent social behavior, with millisecond resolution and accuracies that can out-perform human observers. SimBA does not require specialized video acquisition hardware nor extensive computational background. Standard descriptive statistical analysis, along with graphical region of interest annotation, are provided in addition to predictive classifier generation. To increase ease-of-use for behavioural neuroscientists, we designed SimBA with accessible menus for pre-processing videos, annotating behavioural training datasets, selecting advanced machine learning options, robust classifier validation functions and flexible visualizations tools. This allows for predictive classifier transparency, explainability and tunability prior to, and during, experimental use. We demonstrate that this approach is flexible and robust in both mice and rats by classifying social behaviors that are commonly central to the study of brain function and social motivation. Finally, we provide a library of pose-estimation weights and behavioral predictive classifiers for resident-intruder behaviors in mice and rats. All code and data, together with detailed tutorials and documentation, are available on the [SimBA GitHub repository](#).

Graphical abstract

SimBA graphical interface (GUI) for creating supervised machine learning classifiers of rodent social behavior.

(a) Pre-process videos.

SimBA supports common video pre-processing functions (e.g., cropping, clipping, sampling, format conversion, etc.) that can be performed either on single videos, or as a batch.

(b) Managing pose-estimation data and creating classification projects.

Pose-estimation tracking projects in DeepLabCut and DeepPoseKit can be either imported or created and managed within the SimBA graphical user interface, and the tracking results are imported into SimBA classification projects.

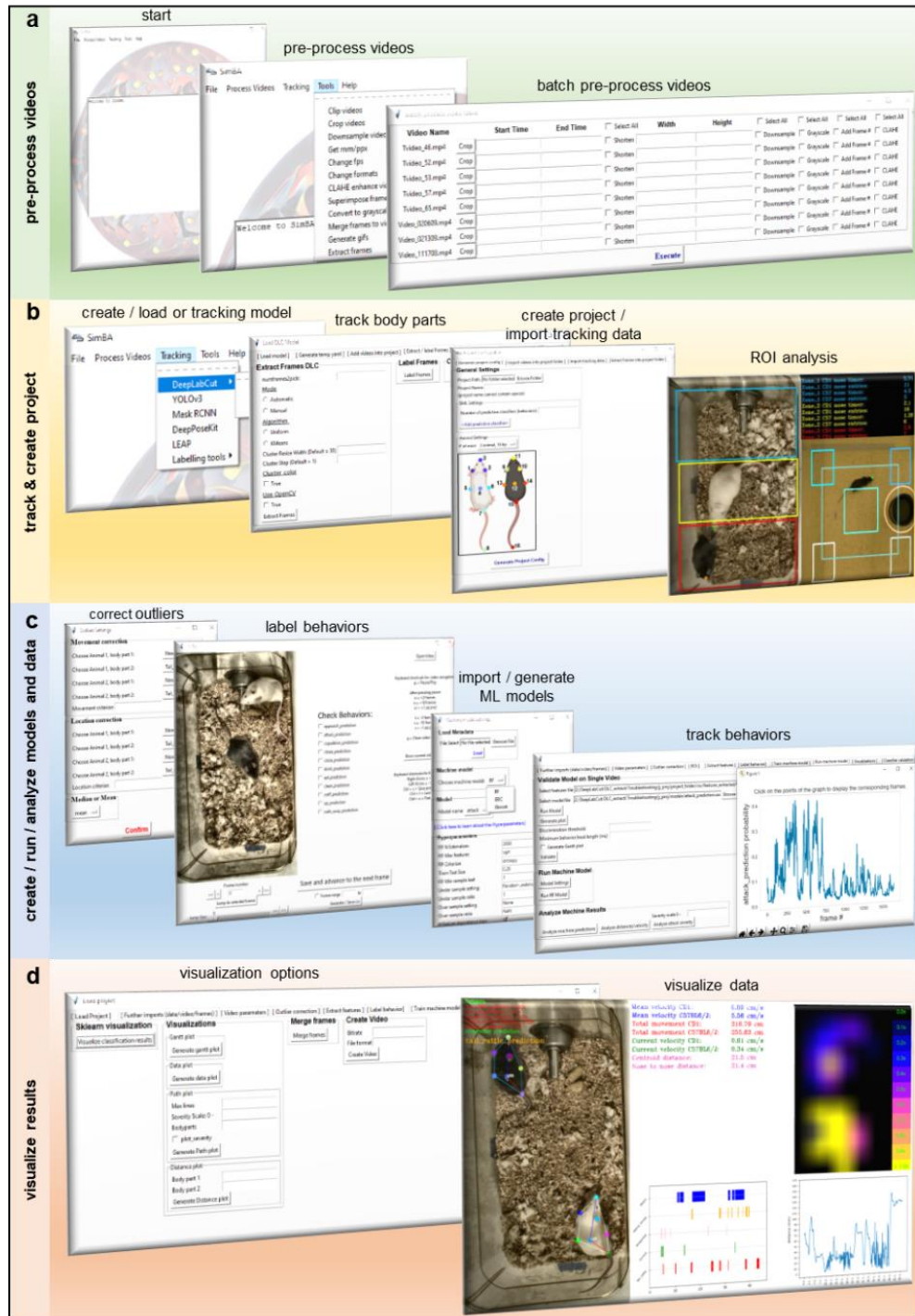
SimBA also supports user-drawn region-of-interests (ROIs) for descriptive statistics of animal movements, or as features in machine learning classification projects.

(c) Create classifiers, perform classifications, and analyze classification data.

SimBA has graphical tools for correcting pose-estimation tracking inaccuracies when multiple subjects are within a single frame, annotating behavioral events from videos, and optimizing machine learning hyperparameters and discrimination thresholds. A number of validation checkpoints and logs are included for increased classifier explainability and tunability prior to, and during, experimental use. Both detailed and summary data are provided at the end of classifier analysis. SimBA accepts behavioral annotations generated elsewhere (such as through JWatcher) that can be imported into SimBA classification projects.

(d) Visualize classification results. SimBA has several options for visualizing machine learning classifications, animal movements and ROI data, and analyzing the durations and frequencies of classified behaviors.

See the [SimBA GitHub repository](#) for a comprehensive documentation and user tutorials.



Introduction

Maladaptive social behaviours are theorised as both a cause and consequence of many neuropsychiatric disorders including depression^{1,2}, substance abuse³ and PTSD⁴⁻⁷. Such social impairments can result from compromised reward evaluations where the motivation for maintaining healthy relationships are persistently diminished⁸. As such, the treatment of maladaptive social behaviour presents an important strategy for improving mental health⁹⁻¹². However, despite the established role of social behaviour in psychiatric disorders and treatments, it remains unclear how social context interact with biological functions to produce protracted and chronic sequelae¹⁰. These insights depend on experimental animal model systems where detailed behavioral ethograms can be combined with precise manipulations of brain functions to reveal biological causes¹²⁻¹⁴.

Unfortunately, the preclinical study of complex social behavior is impeded by the difficulties associated with reliably scoring complex social interactions. Analyses are typically performed by a trained individual, or preferably several trained individuals for appropriate inter-rater-reliability validation, observing social behaviors and manually scoring previously defined events and their durations. The approach is arduous, non-standardized and susceptible to confounds produced by observer drift, long analysis times, and poor inter-rater-reliability. These confounds often prevent the detailed study of complex social repertoires in larger datasets, and notably provide far lower temporal resolution than most modern methodologies such as *in vivo* electrophysiological recording, fiber photometry recording and single-cell calcium endomicroscopy¹⁵⁻¹⁸. These important omissions may contribute to our limited biological understanding of maladaptive social behavior and their neurobiological underpinnings.

To overcome this, a range of elegant open-source tools (Table 1) that use various forms of computer vision with synchronized RFID-tracking data and/or depth camera or multi-camera 3D systems have been developed for automated and precise classifications of animal behaviour¹⁹⁻²⁴. Such methods can permit online classifications in semi-natural environments and are a foundation for impending closed-loop monitoring and forecasting systems²⁵ in behavioural neuroscience, but do require significant investment in specialized hardware and a working knowledge of computer science approaches. Parallel advances in animal tracking have produced accessible open-source pose-estimation tools for accurate tracking of experimenter-defined body-parts in noisy and variable environments²⁶⁻²⁹ (Table 1). These open-source initiatives have proven to be both far less expensive to execute, and provide better animal tracking outcomes, than currently available commercial products³⁰. The collective developments promise decreased workload and reduced experimenter interference while introducing sampling frequencies and scoring accuracies that are compatible with other modern neuroscience techniques. Automated classification techniques may also reduce confounds and increase replicability through computer models based on standardized cross-laboratory definitions of behavioral repertoires, made available in libraries for the scientific community. Together, these approaches may solidify the study of rodent social behavioral under the umbrella of the recently-reinvigorated field of computational neuroethology³¹⁻³⁶ and increase our understanding of naturalistic unrestrained behavior within the context of neural function.

However - despite these developments - manual experimenter annotations persist as the conventional approach for studying social behavior in experimental animals. We have identified a number of interrelated obstacles that prevent widespread implementation of machine learning techniques for behavioral neuroscience, and we are presenting this perspective in an upcoming in-depth review³⁷. Now briefly, and most importantly, these include (i) the moderate to advanced computational/engineering skills and significant time investments that are necessary to implement most machine learning approaches, and (ii) the “black-box” nature and lack of explainability^{38,39} and validation that current approaches are often paired with. Put simply, the learning curves for novice users are steep and often dependent on skill-sets not commonly taught within behavioral neuroscience curricula, and the interpretation of their outcomes can be ethereal.

Here we present an open-source method and graphical user-interface (GUI) called SimBA (**S**imple **B**ehavioral **A**nalysis; Fig. 1) that is used to generate supervised decision ensembles of complex social behaviors from basic video recordings of mouse and rat dyadic encounters. SimBA uses features generated from pose-estimation tracking data together with experimenter-made annotations to generate random forests algorithms that accurately classify behavioral patterns in experimental videos. We package SimBA with a range of accessible tools for processing video data and advanced functions for validating and evaluating predictive classifiers and visualizing machine classifications. SimBA has been developed for analysing complex social interactions but has been adopted for a variety of non-social behavioral protocols at other labs and institutes. We show that the method is flexible by accurately predicting an extensive set of behaviours relevant for studying the neural mechanisms

underlying social motivations in mice and rats. All methods, code, and data together with detailed tutorials are available on the [SimBA GitHub repository](#).

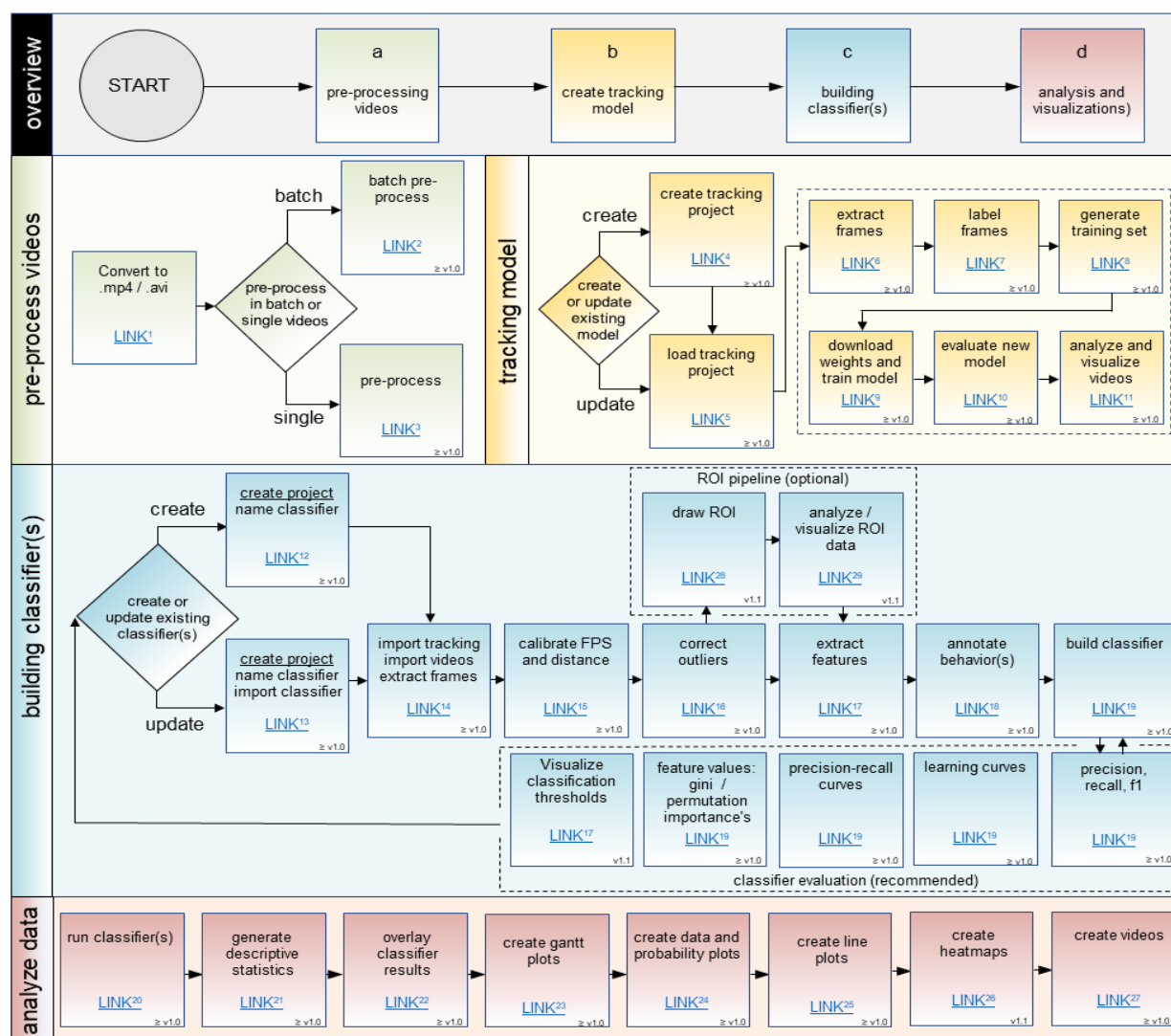


Figure 1. Flowchart depicting the SimBA workflow for creating supervised machine learning classifiers.

An active-link version of this figure is found on the [SimBA GitHub repository](#). (a) *Pre-process videos*. SimBA provided built-in pre-processing options (trim, crop, enhance, down-sample) that can be applied to either single or multiple videos prior to pose-estimation body-part tracking. Videos are CLAHE (Contrast Limited Adaptive Histogram Equalization) enhanced if recorded in greyscale and/or deficient resolution. (b) *Animal tracking*. The pre-processed videos are analyzed using the appropriate DeepLabCut/DeepPoseKit pose-estimation model. Pre-trained pose-estimation model weights for a variety of experimental protocols are available to download from the [SimBA OSF repository](#). (c) *Building classifiers*. Gross inaccuracies in pose-estimation tracking are corrected, and machine learning features (e.g., movements, distances etc.) are calculated from the corrected tracking data. The videos are annotated for the behaviors of interest using the SimBA event-logger, and the annotated events are concatenated with the corrected pose-estimation tracking data. Random forest hyperparameters and discrimination thresholds are tuned, and classifier performances are evaluated. Our random forest classifiers for resident-intruder protocols can be imported into SimBA projects and are available to download from the [SimBA OSF repository](#). (d) *Analysis and visualizations*. The patterns of the classified behaviors, and critical machine learning features (velocities, total movements etc.), are analyzed and visualized. See [SimBA on YouTube](#) for visualization examples.

Methods

SimBA workflow overview

A flowchart depicting the workflow of SimBA is shown in Fig. 1, with links to access online tutorials for the specific SimBA modules. An active-link version of Fig. 1 is found on the [SimBA GitHub repository](#).

Hardware requirements. The recommended system requirements and detailed installation instructions can be found on the [SimBA GitHub repository](#). Briefly, the minimum specifications are an i7 CPU, 16GB RAM, and SSD hard drive. The recommended specifications are an i9 CPU, 32GB RAM, an SSD hard drive, and NVIDIA RTX2080Ti GPU. Animal pose-estimation tracking (i.e., DeepLabCut²⁷ / DeepPoseKit²⁶), component of the SimBA pipeline, requires GPU support. Importantly, SimBA can accept imported pose-estimation data that has been previously generated or generated on a separate system. The visualization steps of SimBA rely on basic functions in the OpenCV and FFmpeg libraries, with are free software suits for video editing and computer vision, and benefit from higher-end CPUs and SSDs.

Installation options. Two supported versions of SimBA (Fig. 2) are available to download from the SimBA GitHub repository: *SimBAxTF* and *SimBA*. The *SimBAxTF* version requires TensorFlow⁴⁰, local GPU-support, and has integrated graphical menus for creating convolutional neural networks through the DeepLabCut²⁷ and DeepPoseKit²⁶ packages. Conversely, the self-contained SimBA version does not require TensorFlow, or local GPU-support, and accepts imported pose-estimation tracking data generated separately within the DeepLabCut or DeepPoseKit notebooks and interfaces.

Software. SimBA is written for Windows compatibility using only open-source software. The code was written in python³⁴¹. The GUI was written in tkinter⁴². The GUI uses DeepLabCut²⁷ or DeepPoseKit²⁶ for body-part labeling and body-part pose-estimation. The SimBA code for generating machine learning behavior classifiers and visualizations rely primarily on scikit-learn⁴³, OpenCV⁴⁴, FFmpeg⁴⁵, and imblearn⁴⁶. A list of SimBA package dependencies and detailed installation instructions can be found on the [SimBA GitHub repository](#).

Datasets. We used SimBA to create social predictive classifiers from three experimental protocols (Table 2): mouse resident-intruder, rat resident-intruder, and the [Caltech Resident-Intruder Mouse dataset \(CRIM13\)](#)^{19,47}.

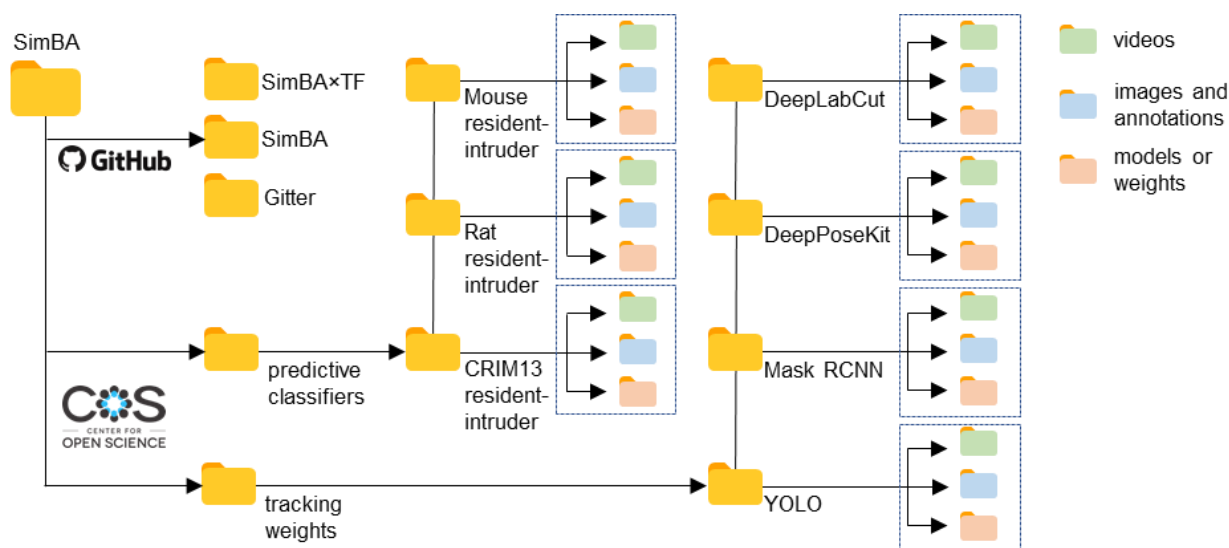


Figure 2. The SimBA open-source code-base and folder structure of the open-access datasets for creating social behavior predictive classifiers. Two supported versions of SimBA are available to download from the [SimBA GitHub repository](#); SimBA with (SimBAxTF) and without (SimBA) integrated TensorFlow-support for convolutional neural networks and pose-estimation animal tracking through DeepLabCut and DeepPoseKit. Support is available through the [SimBA Gitter](#) instant messaging service and chat room. Go to the [SimBA Open Science Framework \(OSF\) repository](#) to download the annotations, videos, random forest models, and tracking weights within predictive classification projects. The [SimBA OSF repository](#) also contains image annotations and network weights for tracking white and black coat-colored animals using alternative neural network architectures (e.g., mask RCNN and YOLO).

Mouse and rat resident-intruder videos were recorded and annotated at the University of Washington and Washington State University, respectively. CRIM13 is an extensive open-access online catalogue of resident-intruder videos accompanied by detailed annotations of different social and non-social behaviors. The mouse and rat resident-intruder datasets are available to download from the [SimBA OSF repository](#). See the [SimBA GitHub repository](#) or the [SimBA OSF repository](#) for a detailed depiction of the folder structure and data that is available for download (Fig. 2).

Video pre-processing. Video recordings were pre-processed using tools available in SimBA. We shortened, cropped, and saved videos and frames in RGB, greyscale and CLAHE (Contrast-limited adaptive histogram equalization)⁴⁸ enhanced formats at variable frame rates. The CRIM13 dataset was recorded in SEQ file format and converted to MP4 format using SimBA. We also used SimBA to extract frames at specific time-periods, down-sample video resolutions, generate gifs and videos. An exhaustive list of SimBA video pre-processing tools and tutorials is available on the [SimBA GitHub repository](#). We used the SimBA to crop, clip, CLAHE enhance, and convert 232 videos in the CRIM13 database^{19,47}.

Classifier operational definitions. We created a detailed operational definition for each classified behavior. The operational definition described the behavior and its characteristic initiation, duration and end. For the mouse-resident intruder protocol (Table 3), we created operational definitions for 10 predictive classifiers: *attack, pursuit, lateral threat, anogenital sniffing, allogrooming normal, allogrooming vigorous, mounting, scramble, flee, and upright submissive*. For the rat resident-intruder protocol (Table 4), we created operational definitions for 7 predictive classifiers: *attack, anogenital sniffing, lateral threat, approach, boxing, avoidance, and submission*. The CRIM13 dataset was previously annotated for 11 different behaviors: *approach, attack, chase, circle, copulation, drink, eat, sniff, up, clean, and walk away*. For further information, see the [CRIM13 website](#).

Resident-intruder protocol. For the mouse resident-intruder protocol, we recorded dyadic encounters between two male mice (cage size: 28x17x14cm) or two female mice (cage size: 28x19x12cm) in clear polycarbonate home-cages with fresh bedding, or two male mice in Med-Associates operant chambers (chamber size: 21x18x13cm) using protocols detailed elsewhere^{49,50}. The intruder was a black coat-coloured C57BL/6J mouse (Jackson Labs, #000664), and the resident was a white coat-coloured CD-1 mouse (Charles River, #022). For the rat resident-intruder protocol, we recorded resident-intruder dyadic encounters between a male intruder Sprague Dawley rat (Simonsen Laboratories) and male resident Long-Evans rat (Simonsen Laboratories) in clear polycarbonate cages with fresh bedding (cage size: 33x46x19cm) using a previously detailed protocol⁵¹.

Video recordings. Male mice were recorded at 30-80fps with USB3.0 cameras (acA2040-120uc - Basler ace, Basler) using fixed-focal length lenses (Edmund Optics, NJ, 16mm/F1.4) from a 90° angle at variable resolutions (W:1000-1200px, H:1255-2056px) using the pylon camera software (Basler). Male rats were recorded at 60fps and 1280x720 resolution at a 90° angle from above using a Logitech C922 USB web-camera. Importantly, all recorded videos used to build behavioral classifiers were re-sampled in SimBA to 30fps before being used to generate machine learning classifiers. All video recordings can be found on the [SimBA OSF repository](#). The Caltech Resident-Intruder Mouse (CRIM13) dataset was recorded at 25fps and 640x480 resolution from a 90° angle.

Project management. SimBA has separate drop-down toolbar menus for creating and loading animal tracking projects and predictive behavior classification projects. Animal tracking data is generated within animal tracking projects and subsequently exported to predictive classification projects. The tracking data is used to create predictive classifiers or analyzed using previously generated predictive classifiers. Within the classification project menus, the user can also analyze animal movements and behavioral patterns based on [user-drawn region-of-interests \(ROIs\)](#).

Animal pose-estimation tracking. DeepLabCut and DeepPoseKit projects were accessed and managed through the TRACKING>DEEPLABCUT and TRACKING>DEEPOSEKIT toolbars of SimBA. We used the DeepLabCut and DeepPoseKit labeling interfaces to annotate eight body-parts on each of the two animals (Fig. 3a). For alternative protocols, SimBA supports any body labeling schematic if first defined by the user in the CREATE PROJECT menu of SimBA (Fig. 3c-d). All generated tracking models, together with the annotated images, are available for download on the [SimBA OSF repository](#). The [SimBA OSF repository](#) also contains annotations and models for detecting black and white coat-colored mice through alternative convolutional learning architectures (e.g., mask RCNN⁵², YOLO⁵³).

Behavior classifications.

Social classifiers. We created and managed machine classification projects using the FILE> CREATE PROJECT menu and FILE>LOAD PROJECT menu in SimBA. We imported the pose-estimation tracking data in CSV file format together with the accompanying videos into each project. We used SimBA to extract the individual frames for each of the videos in the projects (*note*: extraction of frames is a CPU-intensive and time-consuming process that depend on the frame-rate, length, and resolution of the videos). Creating frames for individual videos, however, is only necessary for annotating behaviors and creating new classifiers in SimBA and is not necessary for visualizing and analyzing videos. See the [SimBA GitHub repository](#) for tutorials and recommended approaches for extracting frames in different use-case scenarios.

Time and distance standardization. We used SimBA to standardize distance (pixel/mm) and time (frames per second) measures across different video recordings. SimBA allows the user to define and draw a known distance in the video (for example, the arena length or the arena width) and this distance is used to convert Euclidean pixel distances to millimeter distances. Furthermore, SimBA automatically registers the frame rate of each video, and this information is used to standardize calculations across rolling time windows.

Correcting inaccurate tracking. We designed two outlier correction tools that identify and correct gross pose-estimation tracking inaccuracies by detecting outliers based on movements and locations of body-parts in relation to the animal body-length (Fig. 4a). To compute an initial reference value, SimBA first calculates the mean Euclidean millimeter distance between two user-defined body-parts across the entire recording (L). We set the body-parts to be the nose and the tail-base. We also set two criterion values, one criterion value for 'movement outliers' (V_1) and one criterion value for 'location outliers' (V_2). A body-part coordinate was first corrected as a 'movement outlier' if the movement in Euclidean pixels of the body-part, across two sequential frames, was equal or more than $L \times V_1$ (Fig. 4b, top). If a body-part was found to be a movement outlier in six consecutive frames a

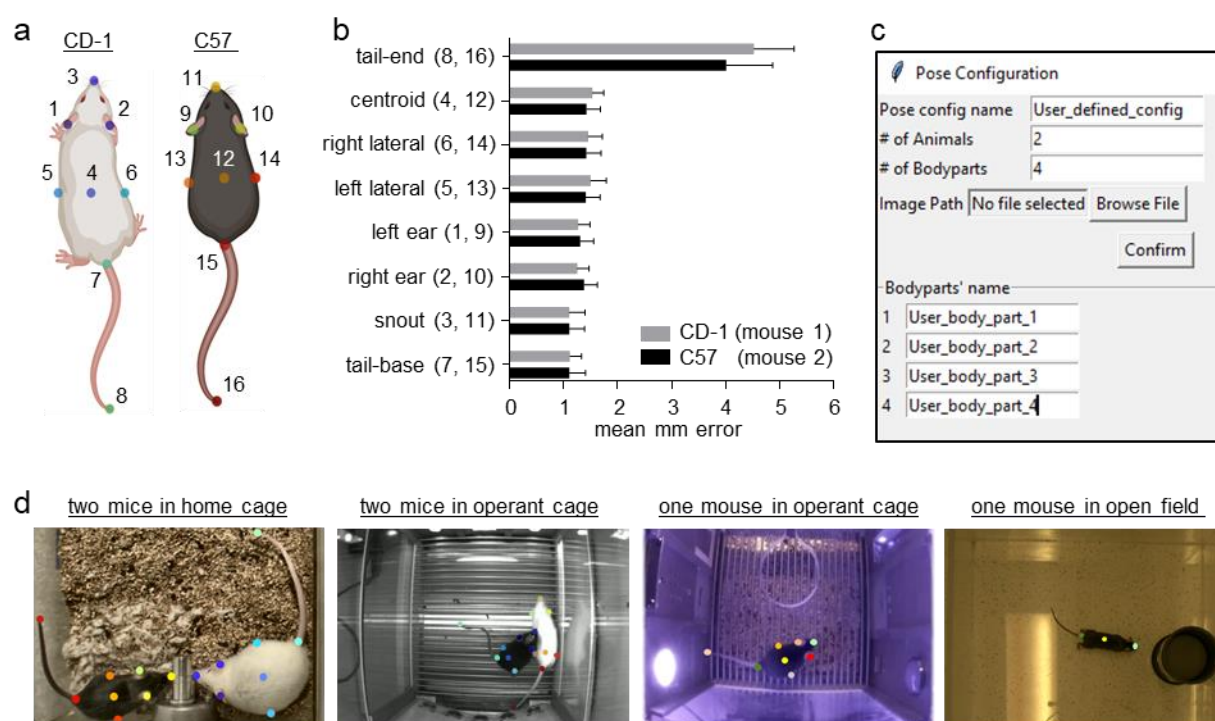


Figure 3. Animal pose-estimation body-part tracking. (a) Schematic representation of the 16 body-parts annotated and tracked in the resident-intruder protocols using DeepLabCut and DeepPoseKit. The adjacent numbers represent the order which the body-parts were annotated in the DeepLabCut and DeepPoseKit annotation interfaces. (b) The mean millimeter tracking errors for the body-parts in the mouse RGB resident-intruder model. (c) The SimBA flexible annotation module for creating and importing user-defined body-part tracking schemas for generation of supervised machine learning classifiers. (d) Examples of body-part tracking schemas created in SimBA for pose-estimation and supervised machine learning classifiers.

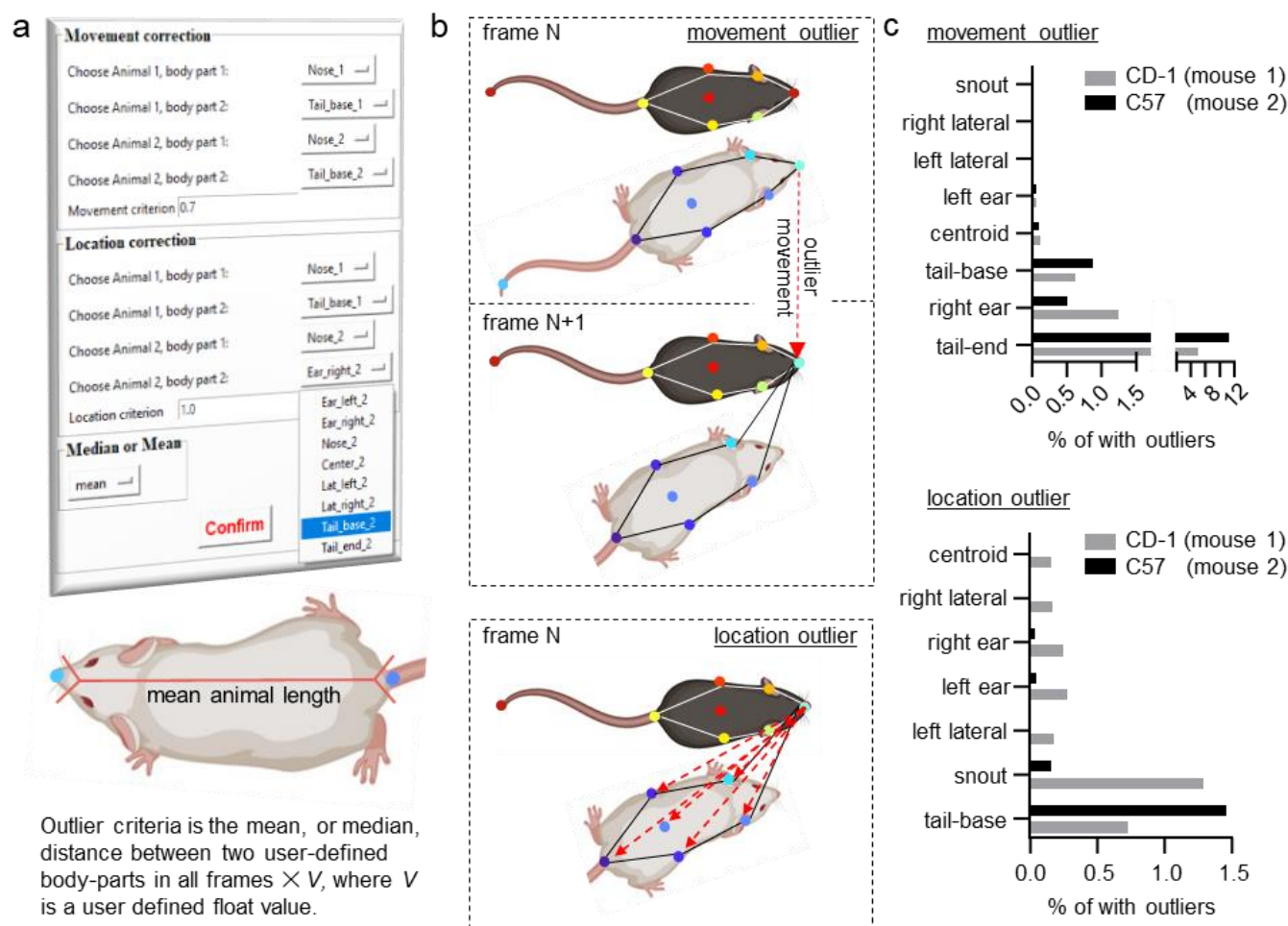


Figure 4. Tools in SimBA for removing inaccuracies of pose-estimation body-part tracking. (a) SimBA calculates the mean or median distance between two user-defined body-parts across the frames of each video. We set the user-defined body-parts to be the nose and the tail-base of each animal. The user also defines a *movement criterion* value, and a *location criterion* value. We set the movement criterion to 0.7, and location criterion to 1.5. Two different outlier criteria are then calculated by SimBA. These criteria are the mean length between the two user-defined body parts in all frames of the video, multiplied by the either user-defined *movement criterion* value or *location criterion* value. SimBA corrects movement outliers prior to correcting location outliers. (b) Schematic representations of a pose-estimation body-part ‘movement outlier’ (top) and a ‘location outlier’ (bottom). A body-part violates the movement outlier criterion when the movement of the body-part across sequential frames is greater than the movement outlier criterion. A body-part violates the location criteria when its distance to more than one other body-part in the animals’ hull (except the tail-end) is greater than the location outlier criterion. Any body part that violates either the movement or location criterion is corrected by placing the body-part at its last reliable coordinate. (c) The ratio of body-part movements (top) and body-part locations (bottom) detected as outliers and corrected by SimBA in the RGB-format mouse resident-intruder data-set. For the outlier corrected in rat and the CRIM13 datasets, see the [SimBA GitHub repository](#).

new ground truth value was sampled. Second, a body-part coordinate was corrected as a ‘location outlier’ if the distance in Euclidian millimeters of that body-part, to at least two other body-parts belonging to the same animal (excluding the tail-end), was equal or more than $L \times \sqrt{2}$ (Fig. 4b, bottom). Body-parts detected as outliers were corrected to their last reliable coordinate. The SimBA outlier correction tools generate logs of the total number of corrected body-parts, and ratio of corrected body-parts, in each video of the project.

Decision tree features. SimBA combines the registered distance (pixels/mm) and time (frames per second) variables for each video with the corrected tracking data to calculate an exhaustive list of distance, movement, angles, areas, and paths metrics and their deviations and rank for individual frames and across rolling windows²⁴.

The features calculated by SimBA, and how many features that are calculated, depend on the body-parts tracked during pose-estimation. When using a 16-body-part tracking schematic on two animals, SimBA extracts a set of 498 different metrics relevant for classifying social and non-social behaviors in the mouse and rat resident datasets. When using other pre-defined body-part configurations, or user-defined body-part configuration, SimBA calculates a reduced and/or generic feature set based on all individual body-parts and their relationships and movements. Lists and descriptions of the different features calculated in different use-case scenarios are available on the [SimBA GitHub repository](#).

Behavior annotations. SimBA has an in-built event logger that displays individual frames alongside the video. Within the event logger interface, the user selects the behavioral events (defined during project creation) that are occurring in the displayed frames, and SimBA concatenates the logged events with the features calculated from the corrected pose-estimation tracking data. We used the SimBA event logger to annotate frames as containing, or not containing, the behaviors of interest in the mouse and rat resident-intruder videos. A tutorial for using the SimBA event logger is available on the [SimBA GitHub repository](#). In many alternative scenarios, the behavioral events may have already been logged through alternative software tools (e.g., JWatcher, Noldus Observer). In these scenarios it is preferable to build predictive classifiers from previously generated event logs without performing further annotations in SimBA. The exact code will depend on the structure of the previously generated event logs. For example, the CRIM13 dataset was annotated using a MATLAB annotation tool⁵⁴. We concatenated the CRIM13 annotations to the CRIM13 tracking data using a [modifiable script](#) that can be downloaded from the SimBA GitHub repository. We request that individuals with already-annotated datasets contact us for developing scripts to easily import annotations into SimBA.

Classification ensembles. Random forest classifiers are intuitive algorithms generated by majority verdicts from decision trees that split the data along feature values to separate distinct classes (e.g., *attack* or *not attack*)⁵⁵. We used SimBA to create random forest classifiers, with one classifier for each of the behaviors of interest. Each classifier contained 2k decision trees. The number of trees was chosen as a large, computationally feasible, amount⁵⁶. Random forest classifiers accept a range of *hyperparameters* that specify how the decision trees should be generated. SimBA accepts a range of different random forest hyperparameter settings, but users unfamiliar with the available parameters can import recommended or previously successful settings based on classifiers of behaviors with similar frequency and salience. Hyperparameter meta files, that can be imported into SimBA and contain the hyperparameters used to generate the current classifiers, can be downloaded from the [SimBA OSF repository](#). We evaluated each classifier by calculating its *precision*, *recall*, and *F1-scores* after 5-fold shuffle cross-validations on 20% of the datasets annotated in the SimBA event logger. Precision was calculated as

$$precision = \frac{true\ positive}{true\ positive + false\ positive}$$

and denotes the proportion of frames correctly classified as displaying the behavior of all the frames classified as containing the behavior. Recall was calculated as

$$recall = \frac{true\ positive}{true\ positive + false\ negatives}$$

and denotes the proportion of all the frames containing the behavior that were correctly classified as containing the behavior. F1 was calculated as

$$F1 = 2 \times \frac{precision \times recall}{precision + recall}$$

and is the harmonic mean of the precision and recall scores.

Random forest classifiers - as other classification techniques - are sensitive to class imbalances. Class imbalances are prevalent in most datasets and occur when observations of the majority class (i.e., observations of the *absence* of the classified behavior) substantially outweigh observations of the minority class (i.e., observations of the *presence* of the classified behavior). For example, in the current mouse resident intruder dataset *flee* events were present in 1% of the annotated frames ([Table 2](#)). In this scenario the random forest decision trees can classify most *flee* observations correctly by disregarding the presence of *flee* events. To prevent this, SimBA supports several class re-balancing methods, including random under-sampling of the

majority class, and over-sampling of the majority class using SMOTE⁵⁷ or SMOTE-EEN⁵⁸. We maximized the F₁-score of the classifiers by random under-sampling the majority class (i.e., observations of the absence of the classified behavior) at ratios up to 1:46. See Table 2 for the random under-sampling ratios used for each classifier.

Classifier evaluations. We used SimBA to generate classifier learning curves. In these learning curves we evaluated F₁-scores after performing 5-fold cross-validations using 1, 25, 50, 75 and 100% of the shuffled data sets to predict the classified behaviors on 20% of the datasets. Learning curves indicate how inclusion of further logged behavioral events affect classifier performance. Furthermore, we used SimBA to generate precision-recall curves⁵⁹, that visualize how classifiers can be titrated to balance the sensitivity versus the specificity of the classifications⁶⁰ through different discrimination thresholds. We also used SimBA to calculate *mean decrease gini impurity* and *permutation importance* scores which gauge the importance of individual features for correctly classifying individual behaviors⁵⁵. *Mean decrease gini impurity score* or *impurity importance* represents the average gain in purity of a decision tree when the feature is included. *Feature permutation importance* represents the loss of predictive power when the feature, and no other feature, is scrambled. Lastly, we evaluated the F₁ scores of each classifier through 5-fold cross-validations after shuffling the behavioral event annotations made by the experimenter in the training set, and testing the classifier on the un-shuffled, correctly annotated behavioral data⁶¹. The final random forest classifiers for mouse and rat resident-intruder protocols, and the CRIM13 dataset, can be downloaded through the SimBA OSF repository accessible through the [SimBA GitHub repository](#).

Procedural runtimes. We used the python TIMEIT module to measure the execution time of six important SimBA scripts on two computers with different hardware specifications (Fig. 5). We also measured the mean time to extract 1k individual image frames from a video recorded at six different resolutions on the two computers. We executed each script was five times and recorded the mean execution time.

RESULTS

Animal tracking.

We used the DeepLabCut²⁷ interface to label 16 body-parts on a male CD-1 and C57BL/6J mouse in 6765 frames from 50 videos in the resident-intruder protocol recorded in RGB format (Fig. 3a). We labeled a further 1550 frames of female mice in the resident-intruder protocol (kindly shared by Emily Newman, Tufts). The total number of frames annotated was 8315. The mean tracking error in millimeters for each of the 16 body-parts in the mouse home-cage resident intruder protocol, recorded in RGB format, is shown in Fig. 3b. For most body-parts the mean

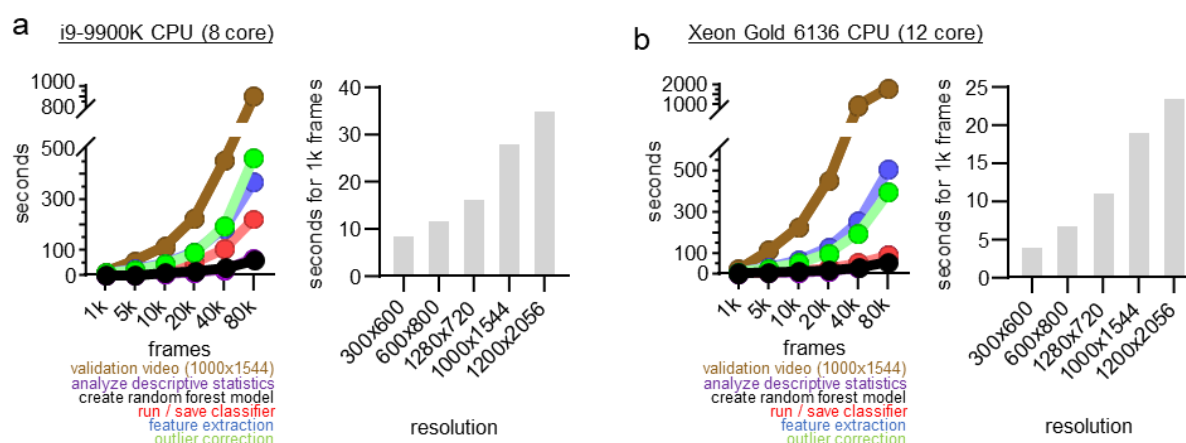


Figure 5. Approximate procedural runtimes for processing different sized data-sets in SimBA. (a) an 8-core Intel i9 CPU and (b) a 12-core Xeon Gold CPU. Time in seconds to perform outlier corrections, feature extraction using 16 body-parts, generating a random forest with 2k trees, performing / saving machine learning classifications, calculating descriptive statistics of machine classifications, generating a validation video, and extracting individual image frames from a video recorded at six different resolutions (videos recorded at 975kbps bitrate). *Note:* only runtimes for creating a validation video and extracting frames depend on the resolution of the videos. See the [SimBA GitHub repository](#) for more information.

Euclidean tracking error was around 1 mm. The tail-ends were associated with larger mean tracking errors between 4-5 mm.

We used the network weights generated from the 8315 annotated images to create additional body-part tracking models for alternative experimental protocols. For these additional tracking models, we labelled a further 800 frames from home-cage environments with male rats, and 3200 CLAHE enhanced images from the Caltech Resident-Intruder Mouse (CRIM13) dataset^{19,47}. All annotations were converted to greyscale and CLAHE enhanced formats, and models were generated based on these annotations. All annotated images, and tracking models, can be downloaded from the [SimBA OSF repository](#).

Procedural Runtime Performance of SimBA functions

The computation speed of several important SimBA functions are shown in Fig. 5. For 1-80k frames on the 8-core Intel i9 CPU-supported computer, pose-estimation outlier correction was performed in 10-465s, feature extraction 5-367s, random forest generation 0-68s, running/saving classifications 4-222s, calculating descriptive statistics 0-67s, and creating a validation video took 11-906s (Fig. 5a). For 1-80k frames on the 12-core Intel Xeon Gold

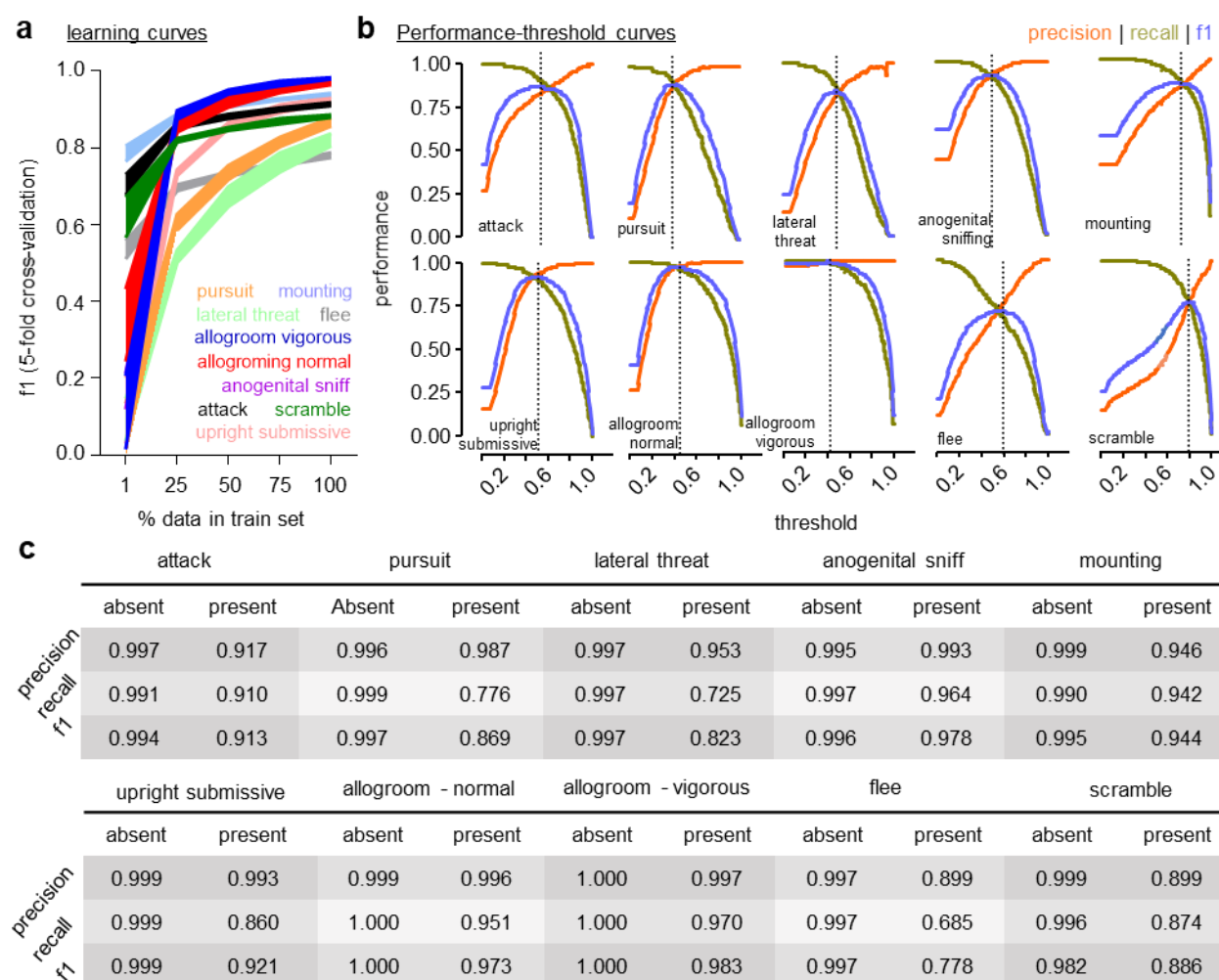


Figure 6. Evaluations of mouse resident-intruder predictive classifiers. (a) Learning curves were created using 2k trees, 5 data splits (1-100%), and with shuffled 5-fold cross-validation at each data split. Errors represent \pm SEM. (b) Classification precision, recall, and F1 scores at different discrimination thresholds. The dotted line represents the discrimination threshold at maximal F1 score. See Table 1 for the optimal thresholds for each classifier. (c) Mean classifier precision, recall, and F1 score evaluated by shuffled 5-fold shuffle cross-validation. See Methods for equations and detailed descriptions of precision, recall, and f1 metrics.

CPU-supported computer, pose-estimation outlier correction was performed in 4-394s, feature extraction 6-504s, random forest generation 0-53s, running/saving classifications 3-90s, calculating descriptive statistics 0-89s, and creating a validation video took 22-1804s (Fig. 5b). Current SimBA efforts are focused on accessible creation of social behavior machine learning classifiers, and future efforts will optimize execution runtimes.

Social Classifiers

Outlier corrections. We used SimBA to correct gross pose-estimation tracking inaccuracies based on impossible locations and movements of animal body-parts (Fig 4a-b). In accordance with the tracking data, the tail-ends represented most of the corrected body-parts based on body-part movements (Fig. 4c). The tail-bases and snouts represented most of the corrected body-parts based on body-part locations. In total, 0.013% of all body-parts were corrected following the movements outlier correction, and a further 0.006% of the body parts were corrected following the location correction. For tracking inaccuracies corrected in other data-sets, see the [SimBA GitHub repository](#).

Classifier datasets. We used the SimBA event logger to annotate video frames as containing, or not containing, the behavior. The number of annotated video frames and the percent of video frames that were annotated as containing the behavior are shown in Table 2. For the classifiers in the mouse resident-intruder dataset, we annotated between 103k and 472k frames. The classified behaviors were annotated as present in 0.73-6.57 % of the video frames. For the predictive classifiers in the rat resident-intruder dataset, we annotated 136k frames and the classified behaviors were annotated as present in 2.48-16.40 % of the frames. Of the original 232 videos in the CRIM13 dataset, we identified 67 videos and 842k frames that contained social-interactions between non-anesthetized black and white coat-colored animals without the presence of a human. We used 64 of the videos and 757k in the training and test sets and saved three videos for validation. The classified behaviors had been annotated as present in 0.36-14.94 % of the video frames.

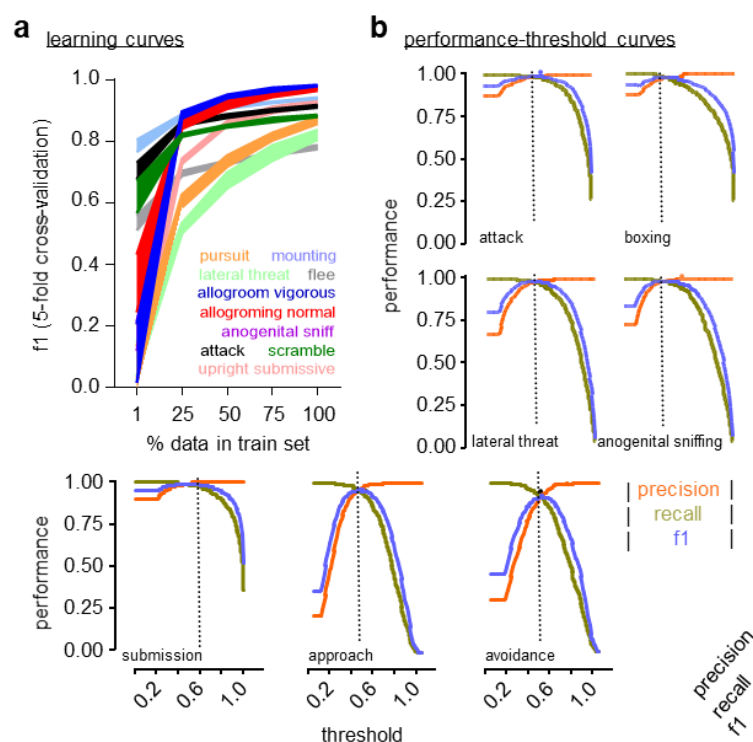


Figure 7. Evaluations of rat resident-intruder predictive classifiers. (a) Learning curves were created using 2k trees, 5 data splits (1-100%) and with shuffled 5-fold cross-validation at each data split. Errors represent \pm SEM. (b) Classification precision, recall, and F1 scores at different discrimination thresholds. The dotted line represents the discrimination threshold at maximal F1 score. See Table 3 for the optimal threshold for each classifier. (c) Mean classifier precision, recall, and F1 score evaluated by shuffled 5-fold shuffle cross-validation. See Method for equations and descriptions of precision, recall, and F1 metrics.

	approach		avoidance	
	absent	present	absent	present
precision	0.994	0.996	0.997	0.990
recall	0.999	0.864	0.998	0.855
f1	0.997	0.925	0.998	0.918

	attack		boxing		lateral threat		anogenital sniff		submission	
	absent	present	Absent	present	absent	present	absent	present	absent	present
precision	0.996	0.991	0.97	0.997	0.994	0.997	0.996	0.992	0.998	0.997
recall	0.997	0.980	1.00	0.937	0.999	0.947	0.999	0.962	0.998	0.987
f1	0.996	0.985	0.998	0.966	0.997	0.971	0.998	0.977	0.998	0.992

Mouse resident-intruder classifier performances. Mean precision, recall and F1-scores for the presence and absence of the behaviors following 5-fold shuffle cross-validation of the final classifiers in the mouse resident-intruder protocol is shown in Fig. 6. Frame-wise classification performance for the *presence* of the behaviors as measured by F1 were 0.778-0.983, precision between 0.899-0.997, and recall between 0.685-0.972. Frame-wise classification performance scores for the *absence* of the behaviors were 0.982 or higher (Fig. 6c). Five-fold cross validation learning curves using 1-100% of the annotated data (Fig. 6a) showed that the number of annotated images positively correlate with F1 score. Precision-recall curves (Fig. 6b) indicated optimal classifier performance as measured by F1 at discrimination thresholds between 0.41-0.79.

Rat resident-intruder classifier performances. Mean precision, recall and F1-scores for the presence and absence of the behaviors following 5-fold shuffle cross-validation of the final classifiers in the mouse resident-intruder protocol is shown in Fig. 7. Frame-wise classification performance for the *presence* of the behaviors as

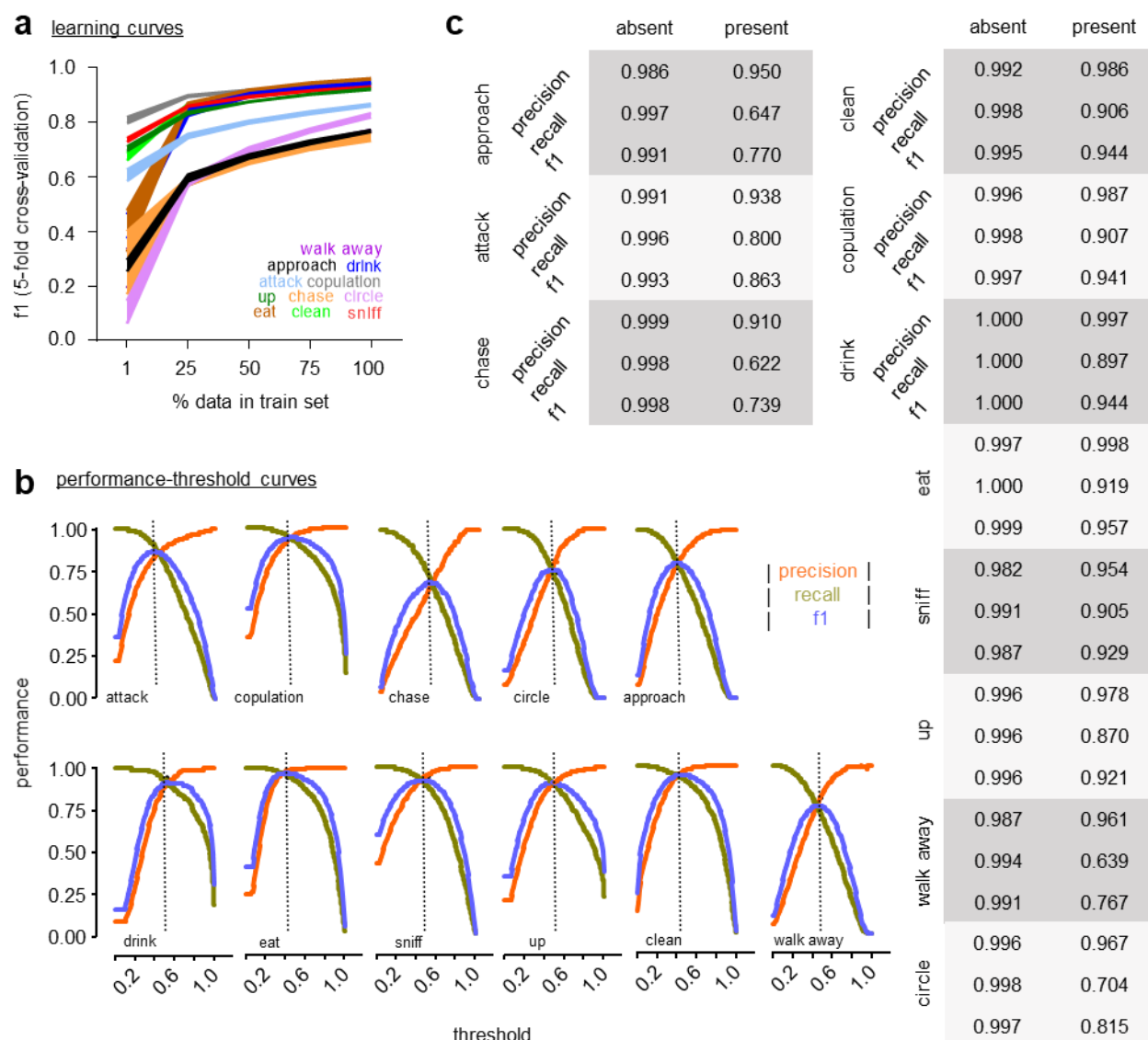


Figure 8. Evaluations of CRIM13 mouse resident-intruder predictive classifiers. (a) Learning curves were created using 2k trees, 5 data splits (1-100%), and with shuffled 5-fold cross-validation at each data split. Errors represent \pm SEM. (b) Classification precision, recall, and F1 scores at different discrimination thresholds. The dotted line represents the discrimination threshold at maximal F1 score. See Table 3 for the optimal threshold for each classifier. (c) Mean classifier precision, recall, and F1 evaluated by shuffled 5-fold shuffle cross-validation. See Method for equations and descriptions of precision, recall, and F1 metrics.

measured by f1 were 0.918-0.985, precision between 0.990-0.997, and recall between 0.855-0.987. Frame-wise classification performance scores for the *absence* of the behaviors were 0.994 or higher (Fig. 7c). Five-fold cross validation learning curves using 1-100% of the annotated data (Fig. 7a) showed that the number of annotated images positively correlate with f1-score. Precision-recall curves (Fig. 7b) indicated optimal classifier performance as measured by f1 at discrimination thresholds between 0.45-0.55.

CRIM resident-intruder classifier performances. Mean precision, recall and F1-scores for the presence and absence of the behaviors following 5-fold cross-validation of the final classifiers in the mouse resident-intruder

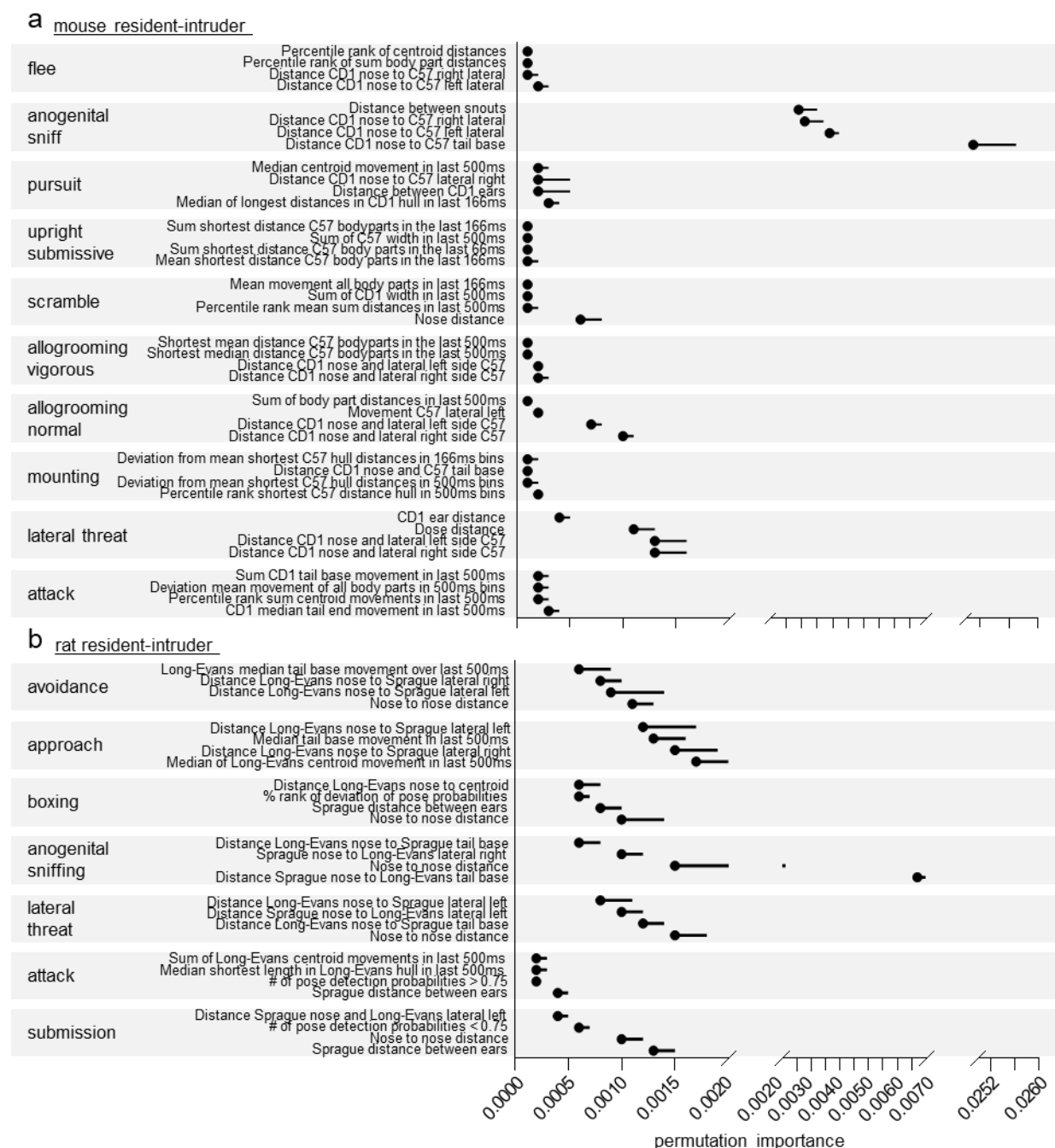


Figure 9. The four most significant features as measured by permutation importances for the classifiers. (a) mouse resident intruder and (b) rat resident-intruder data-sets. Feature permutation importance represents the performance classification degradation when the specific feature, and no other feature, is scrambled. A complete list of feature permutation and gini importance's are available through the [SimBA OSF repository](#).

protocol is shown in Fig. 8. Frame-wise classification performance for the *presence* of the behaviors as measured by F1 were 0.739-0.957, precision between 0.910-0.997, and recall between 0.639-0.919. Frame-wise classification performance scores for the *absence* of the behaviors were 0.991 or higher (Fig. 8c). Five-fold cross validation learning curves using 1-100% of the annotated data (Fig. 8a) showed that the number of annotated images positively correlate with F1-score. Precision-recall curves (Fig. 8b) indicated optimal classifier performance as measured by F1 at discrimination thresholds between 0.40-0.58.

Feature permutation importances. The four most significant features for each classified behavior, as measured by permutation importance's, are shown in Fig. 9. Feature permutation importances represents the performance loss when the specific feature is scrambled. Greater permutation importance values represent a higher feature significance for correct classifications. For complete lists of feature significances for each classified behavior as measured by *mean decrease gini impurity* and permutation importance's, see the [SimBA OSF repository](#). For example, *attacks* in the mouse resident- intruder dataset (Fig. 9a) were classified - in part - based on the amount of movements, the percentile rank of movements / deviation of movements relative to the mean movements in the video. Attack classifications in the rat resident-intruder dataset were also, in part, based on amount of movements, but also the presence of reduced pose-estimation detection probabilities and decreased metric distances between the body-parts in the animal hulls. This can be explained by rats, but not mice, typically assuming stable close-contact *rearing* postures during attack bouts that decrease pixel distances between body-parts and may cause occlusion of individual body-parts.

Annotation scrambling. We performed 5-fold cross-validations where we randomly shuffled the human-made behavior annotations in the training set, before testing the classifier generated from the shuffled data on the correctly annotated, non-shuffled, training set⁶¹. Shuffling the annotations in the training sets to 0.00-0.0036 decreased F1-scores (Fig. 10).

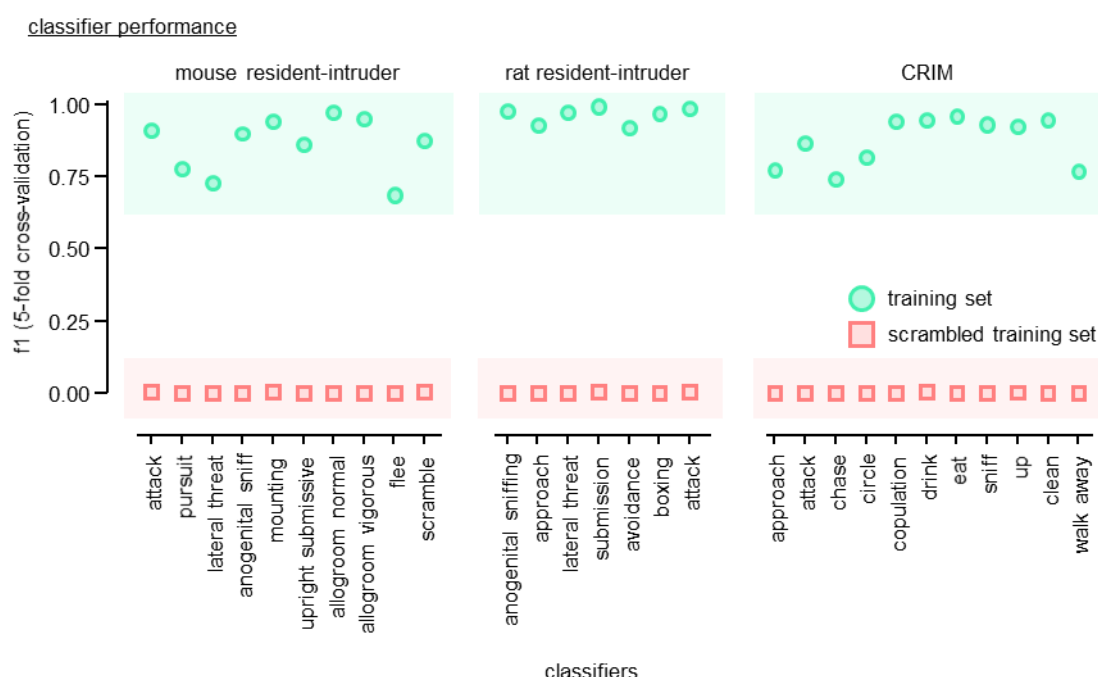


Figure 10. Classifier performance after randomly scrambling the human-made annotations in the training set. Performance was evaluated as F1-score for the presence of the target behavior, measured by shuffled 5-fold cross-fold validation after randomly scrambling the human annotations in the training set. The classifiers were tested using the un-scrambled, correctly annotated, test sets. The green circles represent the performance of the classifiers when trained using un-scrambled annotations. Errors represent \pm SEM (*note*: error bars are present but not discernible in the graph).

Discussion

Open-source machine learning tools for behavioral neuroscience are being released at an intensifying pace³¹. We have identified, and attempted to solve, several issues that prevent a general adoption of machine learning techniques within the field. In many scenarios the required hardware, programming and computational familiarity, and the sensitivity of methods to the experimental conditions, may result in experimenters persisting with manual annotation methods. To advance access to machine learning methods we built SimBA, a software interface that can be used on standard computers by individuals with limited coding experience to implement and generate supervised machine learning algorithms for complex social behavior. We have conscientiously selected a supervised approach, and incorporated a number of explainability and validation checkpoints^{38,39}, to ensure that researchers unfamiliar with these approaches are confident in the accuracy and validity of the classifiers they generate. We have supplied [extensive documentation](#) and [use-case scenarios](#) to simplify the iterative process of tuning supervised predictive classifiers to researcher expectations.

Here we use SimBA to create a battery of 28 predictive classifiers for behavioral repertoires relevant for studying social motivations in rats and mice^{62,63}. We make SimBA, together with the generated classifiers and detailed tutorials, available online to expedite implementation of machine scoring complex behaviors within labs that have limited prior programming experience. Social classifications are readily consolidated with the analysis of calcium indicators or electrophysiological responses and we aim to integrate such alignment options for prevalent neural recording systems within the SimBA graphical interface (Fig. 11).

We make the methods accessible by incorporating video pre-processing options, software support for importing pre-defined machine learning hyperparameters, performing common model evaluation techniques, and tools for visualizing the machine learning classifications. We demonstrate that SimBA is flexible by generating social predictive classifiers across multiple labs for both mice and rats, and by creating accurate classifiers from a third-party dataset available online^{19,47}.

We used SimBA to create classification performance learning curves, where different percentages of the total behavioral annotations are used to create the random forest classifiers (Figs. 5-7). The learning curves show that, for most classifiers, significant improvements can be achieved by adding further experimenter-made behavioral annotations. Many labs working with social behavior already possess video data with accompanying annotations, and this data can be used to both expand the battery of classifiers and sharpen the performance of the classifiers generated here. We have made the datasets used to create our current classifiers available online, and we hope that others will consider further data sharing efforts that increase the scope and accuracy of social machine classifications.

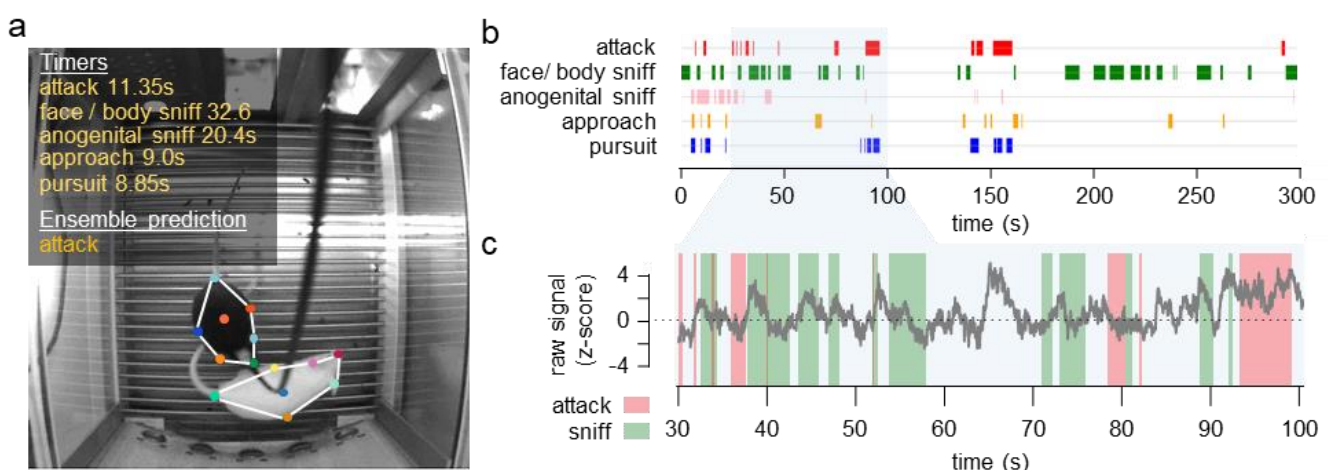


Figure 11. SimBA-generated social predictive classifications aligned with nucleus accumbens (NAc) fiber photometry traces. (a) The mouse resident-intruder protocol within an operant chamber, using a black CD-1 x C57 hybrid resident mouse expressing GRAB_{DA} in the NAc and a white BALB/c intruder mouse. (b) Gantt-plots created in SimBA that display the duration and frequency of classified social behaviors. (c) Z-scored fiber photometry traces (500-550nm) within a 70s-time window. Inset color bars represent classified bouts of attacks (red) and face/body sniffs (green).

Classifier performance improvements can also be achieved through statistical techniques. Accurate prediction of behavior requires handling class imbalances⁶⁴. Datasets in biomedical and behavioral sciences will, in nearly all scenarios, be imbalanced such that observations of the majority class (e.g., observations of the absence of the behavior of interest) will substantially outweigh observations of the minority class (e.g., observations of the presence of the behavior of interest). In our datasets, rare events introduce class imbalances of approximately 1:85 and it is reasonable to assume that alternative behaviors present similar or higher imbalance ratios. SimBA includes a variety of statistical re-balancing tools that address class imbalances^{65–68}. The parameters of these re-balancing tools can be user-defined or imported from similar and previously successfully generated classifiers. Balancing parameters for all 28 classifiers are shown in [Table 1](#), as representative starting points.

The classifiers generated for this manuscript use the DeepLabCut²⁷ default network architectures for pose-estimation positional data, which the subsequent machine learning features are derived from. There are, however, many other effective open-source solutions for animal tracking and pose-estimation^{20,22,69} and SimBA is agnostic to the tools used to extrapolate positional coordinates. SimBA also has an interface for DeepPoseKit²⁶ that permits a range of alternative neural network architectures for pose-estimation that may offer speed and accuracy advantages⁷⁰. Others have also successfully implemented YOLO-based approaches⁷¹. Forthcoming releases of tracking packages promises individual identification of similarly coat-colored animals interacting in groups and we look forward to incorporating such significant developments into SimBA as they are released.

Social interactions - even between differently coat-coloured animals – introduce tracking and pose-estimation challenges that become aggregated by the turbulent interactions and substantial occlusion present in social behavior protocols. For example, DeepLabCut and DeepPoseKit occasionally attribute body-parts to the incorrect animal despite extensive annotation data. We approached the issue post-hoc and designed body-part correction tools that identify and correct movements and distances that are unrealistic based on experimenter-defined criteria. Although the two correction tools eliminate unambiguous outliers, further performance improvements may be obtained by more refined and flexible machine learning methods^{72–74} and/or data cleaning techniques that incorporates variable criteria for the different body-parts. We are currently working on evaluating such techniques and their potential benefits for generating random forest and other decision ensembles for rodent social behavior.

Future directions and challenges

There are some important caveats to be aware of, or account for, during the creation of predictive classifiers with SimBA. Although SimBA can analyze two animals of different coat colors, or a single animal of any coat color, our method does not currently transfer to two or more animals of the same coat color. Current gold-standard methods for performing individual identifications in groups of similarly coat-colored individuals use RFID-data and depth camera systems^{21,22}. Additional advances in machine vision has also addressed identification issues through dual segmentation/identification networks^{75,76} or by including measures of optical flow⁷⁷ and part affinity fields⁷³. Accurate mask-RCNNs⁷⁸ and YOLO-based approaches⁷¹ may also be promising platforms for calculating the rich and relevant feature space that is required for individual identification in social behavior protocols. Identification issues could also be circumvented by adjusting experimental protocols and using differently coat colored mice of the same strain when possible⁵⁰.

SimBA uses comprehensive and generic feature sets (e.g., movements and distances) in random forest designs to permit accurate classifications of variable user-defined target animal behaviors while avoiding critical multi-collinearity issues⁷⁹. In some use-cases, however, random forest designs may not be optimal, and non-specialized feature sets may introduce decision ensemble noise which can negatively affect runtime and the interpretability of feature importance calculations. Commercial alternatives for generating predictive classifiers^{80–82} typically offer exhaustive automated feature engineering/selection methods and evaluate or combine larger batteries of classification techniques when searching for the best alternative. Such tools are also accessible through open-source packages^{83–86} and should be available in future SimBA releases to expedite processing, decrease algorithm complexity, and increase the flexibility of the toolkit. We have nevertheless found random forest designs to be the most robust for classifying the most typically scored aspects of aggression behavior.

A limit to accessibility is that body-part tracking and pose-estimation requires GPU processing. Cloud-based GPU solutions, such as Google Colaboratory / Microsoft Azure / Amazon Web Services, are promoted as alternatives to dedicated GPUs, but this can still be a challenging process. Providing pre-trained network weights only goes so

far in addressing accessibility as users typically must add images representing their environment to the original neural network weights, as well as analyze their own new experimental videos. It is feasible that less complicated classifiers, such as with attack predictions that predominately rely on speed and distance metrics, can be accurately predicted (potentially in real-time) by combination of advanced computational methods such as optical flow⁷⁷ and pretrained YOLO models⁵³.

Conclusions

We demonstrate that SimBA flexibly and simply creates accurate supervised machine learning classifiers using feature sets engineered from open-source pose-estimation tracking data. We demonstrate that the approach is robust by creating social classifiers for both rats and mice across multiple labs, and further by creating classifiers from a previously released public datasets available online¹⁹. We built a GUI that makes our method accessible to experimenters regardless of previous programming experience. This is a step towards generating a standardized toolbox of computational models for scoring complex social behaviors in preclinical neuroscience.

Table 1. Open-source computational packages for animal tracking and/or behavioral classification (not comprehensive)

Name	Validated Species	Multiple animals?	Main output	Behavioral classifiers validated in original publication	Mechanism
Ctrax ⁸⁷	Drosophila	Yes	Behavior	Walk, stop, sharp turn, crabwalk, backup, touch, chase	Computer vision
CADABRA ⁸⁸	Drosophila	Yes	Behavior	Lunging, tussling, wing threat/extension, circling, chasing, copulation	Supervised
Autotyping ⁸⁹	Mice	No	Behavior	Home-cage behaviors: drink, eat, groom, hang, micromovement, rear, rest, walk	Supervised
JAABA ²⁴	Drosophila, mice	Yes	Behavior	Flies: walk, stop, crabwalk, backup, touch, chase, jump, copulation, wing flick/grooming/extension, righting, center pivot, tail pivot. Mice: follow, walk.	Supervised
MiceProfiler ⁹⁰	Mice	Yes, if no obstruction	Behavior	Head-head contact, anogenital contact, side by side, approach, leave, follow, chase	Physics engine
Unnamed ^{*91}	Mice	Yes	Behavior	Approach, walk away, circle, chase, attack, copulation, drink, eat, clean, sniff, rear	Supervised
Unnamed ^{*92}	Rats	Yes	Behavior	Rearing, head-head and head – hip contact, approach, leave, follow, mount, intromission, ejaculation	Physics engine
MotionMapper ⁹³	Flies	Yes	Behavior	Grooming: wing, leg, abdomen, head. Wing waggle, running	Unsupervised
Unnamed ^{*94}	Mice	Yes, if distinct	Behavior	Attack, close investigation, mounting	Supervised
MoSeq ^{*95}	Mice	No	Behavior	Dart, micromovement, pause, rear, walk	Unsupervised
Unnamed ⁹⁶	Mice	Yes	Behavior	Direct contact, nose to nose, anogenital sniffing, following, mating, self-grooming	Unsupervised
DuoMouse ⁹⁷	Mice	Yes	Behavior	Sniffing, following, indifferent	Unsupervised
Stytra ⁹⁸	Zebrafish larvae	No	Closed loop tracking & stimulus admin. (real time)		Computer vision
ToxID / ToxTrac ^{99,100}	Fish, ants, mice	Yes	Individual tracking in groups (real time)		Computer vision
MoST ¹⁰¹	Mice	Yes	Individual tracking in groups		Computer vision
idtracker ^{75,76}	Drosophila, fish, ants, mice	Yes	Individual tracking in groups		Unsupervised
LocoMouse ^{*102}	Mice	No	Kinematics		Supervised
DeepLabCut ²⁷	Drosophila, mice, horses, humans, fish	Expected soon	Pose		Supervised
LEAP ²⁸	Drosophila, mice	sLEAP expected soon	Pose		Supervised
DeepFly3d ^{* 103}	Drosophila, humans	No	Pose		Unsupervised
DeepPoseKit ⁷⁰	Drosophila, locusts, zebras	Yes, if distinct	Pose (real time)		Supervised
Sensory Orientation Software ¹⁰⁴	Drosophila, mice, fish	No	Posture & trajectory (real time)		Computer vision
MOTR ¹⁰⁵	Mice	Yes, if distinct	Trajectory		Computer vision
Mouse Behavior Tracker ¹⁰⁶	Mice	No	Trajectory		Computer vision

Asterisk denotes requirement of multiple cameras or special equipment

Table 2. Predictive classifier datasets

	classifier	# annotated frames	# annotated videos	annotations: % present	annotations: present (hh:mm:ss)	annotations: absent (hh:mm:ss)	under sample ratio	test set: frames present	test set: frames absent	optimal threshold
mouse resident-intruder	allogroom normal	334680	26	1.49	00:02:46	03:03:10	19	983	65953	0.48
	allogroom vigorous	334680	26	0.73	00:01:21	03:04:35	0	494	66442	0.41
	attack	203841	37	4.93	00:05:35	01:47:40	9	1920	38849	0.56
	lateral threat	168738	21	1.49	00:01:24	01:32:21	18	503	33245	0.52
	mounting	470294	31	2.88	00:07:31	04:13:45	4	2639	91420	0.72
	pursuit	103538	33	2.02	00:01:10	00:56:22	20	419	20289	0.43
	upright submissive	272304	20	1.13	00:01:43	02:29:34	16	577	53884	0.54
	flee	395852	22	1.08	00:02:23	03:37:33	28	877	78294	0.60
	scramble	323852	18	2.16	00:03:53	02:56:02	3.9	1347	63424	0.79
	anogenital sniffing	103538	33	6.57	00:03:47	00:53:45	8	1268	19440	0.55
rat resident-intruder	attack	136830	12	16.40	00:12:28	01:03:33	0	4437	32119	0.55
	lateral threat	136830	12	9.09	00:06:55	01:09:06	0	9943	26613	0.55
	anogenital sniffing	136830	12	7.50	00:05:42	01:10:19	7.5	2095	22713	0.45
	submission	136830	12	11.27	00:08:34	01:07:27	5	3074	33482	0.59
	boxing	136830	12	4.50	00:03:25	01:12:36	0	1306	23502	0.47
	approach	136830	12	4.23	00:03:13	01:12:48	8.5	1053	23755	0.50
	avoidance	136830	12	2.48	00:01:53	01:14:08	10	686	24122	0.48
CRIM13 resident-intruder	approach	757350	64	4.13	00:20:51	18:01:39	12	6190	145280	0.44
	attack	757350	64	4.10	00:20:42	18:01:48	16	6266	145204	0.40
	chase	757350	64	0.49	00:02:28	18:20:02	46	769	150701	0.58
	circle	757350	64	1.32	00:06:40	18:15:50	22	1982	149488	0.54
	clean	757350	64	7.99	00:40:20	17:42:10	8	12095	139375	0.53
	copulation	757350	64	4.62	00:23:20	17:59:10	12	7080	144390	0.50
	drink	757350	64	0.36	00:01:49	18:20:41	10	570	150900	0.54
	eat	757350	64	2.54	00:12:49	18:09:41	12	3824	147646	0.43
	sniff	757350	64	14.94	01:15:26	17:07:04	0	22757	128713	0.54
	up	757350	64	3.71	00:18:44	18:03:46	6	5575	145895	0.50
	walk away	757350	64	3.89	00:19:38	18:02:52	8	5897	145573	0.50

Table 3. Behavioral operational classifiers: mouse resident-intruder


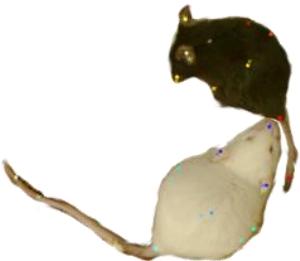


Classifier	Description	Start Frame	Duration of behavior	End Frame
Attack 	Clear physical antagonistic interaction initiated by the Resident mouse. Brief pauses can occur if the Resident mouse remains oriented toward the Intruder.	First frame when the Resident mouse makes physical antagonistic contact with the Intruder. Typically this is characterized by outstretched Resident forepaw(s) contacting the Intruder, while the Resident has an open mouth to initiate a bite. Can also be characterized by the first frame of a slap or quick bite without the forepaws being outstretched.	Short breaks may be present in attack behavior, if Resident is still oriented toward Intruder. Attacks can include tussling, biting, boxing, and corralling as part of the attack bout.	First frame when Resident mouse orients away from Intruder. Typically this is a slight turning of the head to look in a different direction, followed by a relaxation of the body and moving away from the Intruder.
Anogenital sniffing 	Resident mouse is sniffing the anogenital region of the Intruder. Resident must be sniffing at base of tail, not further up on back or on legs.	First frame when the Resident mouse is clearly sniffing anogenital region of Intruder, rather than side, back, or leg.	Uninterrupted sniffing of anogenital region.	First frame when Resident mouse moves head away from anogenital region, either to move away from Intruder or to sniff non-anogenital region.
Lateral threat 	Resident mouse is in proximity (typically less than one body length away from the Intruder) to face of Intruder with back arched and side displayed toward Intruder. Ears are often pinned, with shoulder and side of face nearest to Intruder mouse tilted slightly toward ceiling.	First frame when Resident mouse orients side to Intruder and tilts front of body toward Intruder.	Resident will often circle Intruder or move front half side to side in front of Intruder, feigning attacks prior to actual attack.	First frame when lateral threat posture is dropped. Animal will shift head away to look at other target or will begin an attack. If animal does not attack, the back posture will relax.
Pursuit 	Resident mouse is following in the Intruder's path as the Intruder moves away from the Resident.	First frame when the Resident is moving toward Intruder as Intruder moves away. Typically this is characterized by the Intruder running away after an attack ends and the Resident follows directly after, or when the Intruder walks past the Resident and the Resident markedly changes directions to pursue the Intruder in the Intruder's path.	The Intruder is specifically moving away from the Resident. The Intruder is not sniffing the arena, foraging, etc. Intruder typically moves in a straight line until it reaches the edge of the area, at which point it will turn to face and watch the Resident. Resident is moving along the same path as the Intruder without sniffing or foraging in the arena.	First frame when either mouse stops moving, or the Resident deviates from path of the Intruder.

Table 3 continued. Behavioral operational classifiers: mouse resident-intruder


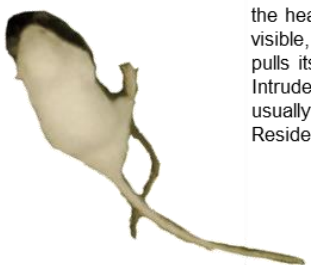

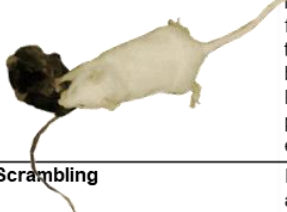





Classifier	Description	Start Frame	Duration of behavior	End Frame
Flee 	Running away from Resident. This is distinct from scrambling because Intruder is actively moving across cage, most typically in a straight line away from Resident.	First frame in which Intruder elongates to move away from the Resident.	Intruder is moving away from Resident in a fast and directed manner.	Intruder deviates to face Resident, alters course in a wandering manner, or slows down or stops.
Mounting 	The Resident is on top of the Intruder. Usually only the head of the Resident is visible, and the Resident pulls its pelvis into the Intruder. The Intruder's back usually arches around the Resident mouse.	The Resident is on top of the Intruder with pelvis is pulled.	The Resident thrusts, or 'humps', while remaining on top of the Intruder with an arched back. The Resident may take quick intermittent breaks while remaining on top of the Intruder. The Resident may be dragged along as the Intruder is moving, but the Resident always returns to thrusting movement on top of the Intruder.	The Intruder pulls away, or the Resident removes itself from the back of the Intruder.
Allogroom normal 	Characterized by Resident placing its paws on the back of Intruder and repeatedly licking its fur; The Intruder is generally occupied with something else.	First frame when Resident places mouth on the Intruder's shoulder/ neck.	Resident constantly licks and bites at fur of Intruder; Intruder stays partially still while also moving head and neck or sniffing resident; Resident moves with the Intruder's movements.	Intruder begins to retreat quickly from the Resident, or the Resident lifts head and retreats.
Upright submissive 	Intruder is standing on hind legs with stomach exposed, forepaws extended or up in the air, and is often leaning backwards away from Resident slightly. Is not pushing or actively engaging with Resident.	First frame in which Intruder is standing on back legs with front paws outstretched and has stomach exposed.	Intruder remains standing on hind legs with forepaws stretched out and stomach exposed. Often is quite still.	Intruder either moves into upright defensive boxing or forepaws touch the ground.
Scrambling 	Intruder is attempting to get away from Resident but is cornered or pinned. Can be in any position (ground, standing, all fours), but this is delineated from other classifiers because Intruder's head is often pointed away from Resident and is clearly scrambling with its paws to try to pull itself away from the Resident.	First frame in which Intruder is attempting to get away from Resident. Different than fleeing because is not yet moving away from Resident and Intruder often has to make several attempts to get around/away from Resident.	Intruder continues to attempt to escape instead of becoming defensive or submissive.	Intruder is either successful and flees, or shifts to other behavior. Often shifts to ground defensive.
Allogroom vigorous 	Characterized by Resident placing its paws on the back of Intruder and repeatedly licking its fur intensely; The Intruder is still and receptive during the grooming.	First frame when Resident places mouth on the Intruder's shoulder or neck.	Resident constantly and aggressively licks and bites at fur of Intruder; Intruder stays still and does not try to move head or neck; behavior is extended.	Resident lifts head and retreats away from Intruder.

Table 4. Behavioral operational classifiers: rat resident-intruder

Classifier	Description	Start Frame	Duration of behavior	End Frame
Attack	Clear physical antagonistic interaction initiated by the Resident rat. Pauses in movement can occur if the Resident remains oriented or moving toward the Intruder.	First frame when the Resident rat makes movement towards the Intruder that ends in a physical attack. A physical attack is typically characterized by outstretched Resident forepaw(s) contacting the Intruder, while the Resident has an open mouth to initiate a bite. Can also be characterized by a quick threatening movement resulting in the rearing up of both rats.	Attacks can include tussling, biting, pinning, boxing, and corralling as part of the attack bout. Breaks in movement may be present in attack behavior. This typically occurs after boxing where both rats are reared up and Resident is oriented towards the Intruder. Breaks in movement can also occur when the Resident has the Intruder pinned in a submissive posture.	First frame when Resident rat orients away from Intruder. Typically this is a slight turning of the head to look in a different direction, followed by a relaxation of the body and moving away from the Intruder.
Submission	The Intruder rat is in a supine posture and is pinned down by the Resident rat. The Intruder can be pinned in a corner or against a wall, but the back of the Intruder close-to horizontal with the bottom of the cage.	First frame when the Intruder rat assumes a supine posture and continuous to be pinned down for at least 1 second.	Struggling usually occurs at the beginning of the submissive posture and is typically followed by breaks in movement with the Resident rat still on top of the Intruder. More bouts of struggling typically happen, and in some cases, may result in the moving of the interaction to a different location during which a submissive posture is still assumed by the Intruder.	First frame when the Intruder rat is no longer in a supine position. Typically characterized by moving upright onto its hind paws or on its side.
Lateral threat	Resident rat is in proximity (typically less than one body length away from the Intruder) to face of Intruder with back arched and side displayed toward Intruder. Ears are often pinned, with shoulder and side of face nearest to Intruder rat tilted slightly toward ceiling.	First frame when Resident rat initiates the movement to orient towards the side of the Intruder, arches back, and tilts front of body toward Intruder.	Resident will often move toward the Intruder or move front half side to side in front of Intruder, feigning attacks prior to actual attack. Can also occur without initiation of an attack upon approach of the Intruder rat as a defensive behavior.	First frame when lateral threat posture is dropped. Animal will shift head away to look at other target or will begin an attack. If animal does not attack, the back posture will relax. Can occur by the Intruder rat increasing proximity to Resident.
Anogenital sniffing	Either rat is sniffing the anogenital region of the other. The rat must be sniffing around the base of tail rather than the back or legs.	First frame when either rat is clearly sniffing the other's anogenital region near the base of the tail. In addition to what is pictured, anogenital sniffing can occur when one rat approaches from the front or side and crawls under the other to investigate.	Uninterrupted sniffing of anogenital region.	First frame when the rat moves head away from anogenital region, usually either to move away from Intruder or sniff non-anogenital region.

Table 4 continued. Behavioral operational classifiers: rat resident-intruder

Classifier	Description	Start Frame	Duration of behavior	End Frame
Boxing 	The Resident and Intruder rats stand on their hind legs in close proximity while pushing and/or grabbing at each other with their front limbs.	First frame when both rats are completely reared up with the hind limbs supporting their weight (as opposed to pushed up against the side of the arena).	A bout of boxing must include contact between the upper, limbs of the rats, whether its pushing and grabbing or light physical contact. This typically occurs within or at the end of an attack bout.	First frame when either rat is no longer fully upright or one rat initiates a movement to return to walking on four limbs, usually moving away from the other.
Approach 	The Intruder rat moves towards the Resident rat.	First frame when the Intruder initiates a movement towards the Resident, or changes direction towards the resident while in the middle of a movement.	The Intruder may be moving towards the Resident slowly or quickly. There may be pauses in movement if the Intruder is moving cautiously with its head directed at the Resident.	First frame when the Intruder has either stopped moving towards the Resident, or makes physical/olfactory contact with the Resident.
Avoidance 	The Intruder rat moves away from the Resident rat.	First frame when the Intruder initiates a movement away from the current or anticipated position of the Resident. Often follows lateral threat, submissive, or boxing behavior.	The Intruder is moving away from the Resident in a continuous movement.	First frame when the Intruder ceases its movement away from the Resident, is attacked, or reaches maximal distance from the Resident (in the case of a long circling movement).

References

1. Martin, L. A., Neighbors, H. W. & Griffith, D. M. The Experience of Symptoms of Depression in Men vs Women: Analysis of the National Comorbidity Survey Replication. *JAMA Psychiatry* **70**, 1100 (2013).
2. Dutton, D. G. & Karakanta, C. Depression as a risk marker for aggression: A critical review. *Aggression and Violent Behavior* **18**, 310–319 (2013).
3. Chermack, S. T., Fuller, B. E. & Blow, F. C. Predictors of expressed partner and non-partner violence among patients in substance abuse treatment. *Drug and Alcohol Dependence* **58**, 43–54 (2000).
4. Chermack, S. T., Fuller, B. E. & Blow, F. C. Predictors of expressed partner and non-partner violence among patients in substance abuse treatment. *Drug and Alcohol Dependence* **58**, 43–54 (2000).
5. Lisak, D. & Miller, P. M. Childhood trauma, posttraumatic stress disorder, substance abuse, and violence. in *Trauma and substance abuse: Causes, consequences, and treatment of comorbid disorders* 73–88 (American Psychological Association, 2003).
6. Dutton, D. G. & Karakanta, C. Depression as a risk marker for aggression: A critical review. *Aggression and Violent Behavior* **18**, 310–319 (2013).
7. Taft, C. T., Creech, S. K. & Murphy, C. M. Anger and aggression in PTSD. *Current Opinion in Psychology* **14**, 67–71 (2017).
8. Bakken, K., Landheim, A. S. & Vaglum, P. Axis I and II disorders as long-term predictors of mental distress: a six-year prospective follow-up of substance-dependent patients. *BMC Psychiatry* **7**, 29 (2007).
9. Abbott, P. J. A Review of the Community Reinforcement Approach in the Treatment of Opioid Dependence. *Journal of Psychoactive Drugs* **41**, 379–385 (2009).
10. Volkow, N. D., Baler, R. D. & Goldstein, R. Z. Addiction: Pulling at the Neural Threads of Social Behaviors. *Neuron* **69**, 599–602 (2011).
11. Elbogen, E. B. *et al.* Protective mechanisms and prevention of violence and aggression in veterans. *Psychological Services* **11**, 220–228 (2014).
12. Heilig, M., Epstein, D. H., Nader, M. A. & Shaham, Y. Time to connect: bringing social context into addiction neuroscience. *Nat Rev Neurosci* **17**, 592–599 (2016).
13. Venniro, M. *et al.* Volitional social interaction prevents drug addiction in rat models. *Nat Neurosci* **21**, 1520–1529 (2018).
14. Venniro, M. & Golden, S. A. Taking action: empathy and social interaction in rats. *Neuropsychopharmacol.* (2020) doi:10.1038/s41386-019-0596-0.
15. Falkner, A. L., Grosenick, L., Davidson, T. J., Deisseroth, K. & Lin, D. Hypothalamic control of male aggression-seeking behavior. *Nat Neurosci* **19**, 596–604 (2016).
16. Ferenczi, E. A. *et al.* Prefrontal cortical regulation of brainwide circuit dynamics and reward-related behavior. *Science* **351**, aac9698–aac9698 (2016).
17. Kim, Y. *et al.* Mapping Social Behavior-Induced Brain Activation at Cellular Resolution in the Mouse. *Cell Reports* **10**, 292–305 (2015).
18. Gunaydin, L. A. *et al.* Natural Neural Projection Dynamics Underlying Social Behavior. *Cell* **157**, 1535–1551 (2014).
19. Burgos-Artizzu, X. P., Dollar, P., Dayu Lin, Anderson, D. J. & Perona, P. Social behavior recognition in continuous video. in *2012 IEEE Conference on Computer Vision and Pattern Recognition* 1322–1329 (IEEE, 2012). doi:10.1109/CVPR.2012.6247817.
20. Hong, W. *et al.* Automated measurement of mouse social behaviors using depth sensing, video tracking, and machine learning. *Proc Natl Acad Sci USA* **112**, E5351–E5360 (2015).
21. Peleh, T., Bai, X., Kas, M. J. H. & Hengerer, B. RFID-supported video tracking for automated analysis of social behaviour in groups of mice. *Journal of Neuroscience Methods* **325**, 108323 (2019).
22. de Chaumont, F. *et al.* Real-time analysis of the behaviour of groups of mice via a depth-sensing camera and machine learning. *Nat Biomed Eng* **3**, 930–942 (2019).
23. Gerós, A., Magalhães, A. & Aguiar, P. Improved 3D tracking and automated classification of rodents' behavioral activity using depth-sensing cameras. *Behav Res Methods* (2020) doi:10.3758/s13428-020-01381-9.
24. Kabra, M., Robie, A. A., Rivera-Alba, M., Branson, S. & Branson, K. JAABA: interactive machine learning for automatic annotation of animal behavior. *Nat Methods* **10**, 64–67 (2013).
25. Schweihoff, J. F. *et al.* DeepLabStream: Closing the loop using deep learning-based markerless, real-time posture detection. *bioRxiv* 2019.12.20.884478 (2019) doi:10.1101/2019.12.20.884478.

26. Graving, J. M. *et al.* DeepPoseKit, a software toolkit for fast and robust animal pose estimation using deep learning. *eLife* **8**:e47994 (2019).
27. Mathis, A. *et al.* DeepLabCut: markerless pose estimation of user-defined body parts with deep learning. *Nat Neurosci* **21**, 1281–1289 (2018).
28. Pereira, T. D. *et al.* Fast animal pose estimation using deep neural networks. *Nat Methods* **16**, 117–125 (2019).
29. Geuther, B. Q. *et al.* Robust mouse tracking in complex environments using neural networks. *Communications Biology* **2**, 1–11 (2019).
30. Sturman, O. *et al.* Deep learning based behavioral analysis enables high precision rodent tracking and is capable of outperforming commercial solutions. *bioRxiv* 2020.01.21.913624 (2020) doi:10.1101/2020.01.21.913624.
31. Datta, S. R., Anderson, D. J., Branson, K., Perona, P. & Leifer, A. Computational neuroethology: a call to action. *Neuron* **104**, 11–24 (2019).
32. Zilkha, N., Sofer, Y., Beny, Y. & Kimchi, T. From classic ethology to modern neuroethology: overcoming the three biases in social behavior research. *Current Opinion in Neurobiology* **38**, 96–108 (2016).
33. Graña, M. & de Lope Asiain, J. A Short Review of Some Aspects of Computational Neuroethology. in *Understanding the Brain Function and Emotions* (eds. Ferrández Vicente, J. M., Álvarez-Sánchez, J. R., de la Paz López, F., Toledo Moreo, J. & Adeli, H.) 275–283 (Springer International Publishing, 2019). doi:10.1007/978-3-030-19591-5_28.
34. Meyer, J.-A. & Wilson, S. W. Computational Neuroethology: A Provisional Manifesto. in *From Animals to Animats: Proceedings of the First International Conference on Simulation of Adaptive Behavior* 29–39 (MITP, 1991).
35. Achacoso, T. B. & Yamamoto, W. S. Artificial ethology and computational neuroethology: a scientific discipline and its subset by sharpening and extending the definition of artificial intelligence. *Perspect. Biol. Med.* **33**, 379–389 (1990).
36. Beer, R. D. *Intelligence as adaptive behavior: an experiment in computational neuroethology*. (Academic Press Professional, Inc., 1990).
37. Goodwin, N., Nilsson, S. & Golden, S. Rage Against the Machine: Advancing the study of aggression ethology via machine learning. *In revision*.
38. Roscher, R., Bohn, B., Duarte, M. F. & Garcke, J. Explainable Machine Learning for Scientific Insights and Discoveries. *IEEE Access* **8**, 42200–42216 (2020).
39. Lundberg, S. M. *et al.* From local explanations to global understanding with explainable AI for trees. *Nature Machine Intelligence* **2**, 56–67 (2020).
40. Abadi, M. *et al.* TensorFlow: Large-Scale Machine Learning on Heterogeneous Distributed Systems. *arXiv:1603.04467 [cs]* (2016).
41. Van Rossum, G. & Drake Jr, F. L. *Python tutorial*. (Centrum voor Wiskunde en Informatica Amsterdam, The Netherlands, 1995).
42. Lundh, F. An introduction to tkinter. URL: www.pythonware.com/library/tkinter/introduction/index.htm (1999).
43. Pedregosa, F. *et al.* Scikit-learn: Machine learning in Python. *Journal of machine learning research* **12**, 2825–2830 (2011).
44. Bradski, G. The OpenCV Library. *Dr. Dobbs's J. Software Tools* **25**, 120–125 (2000).
45. FFmpeg developers. *ffmpeg version N-93294-g056a2ac168*. (2006).
46. Lemaître, G., Nogueira, F. & Aridas, C. K. Imbalanced-learn: A python toolbox to tackle the curse of imbalanced datasets in machine learning. *The Journal of Machine Learning Research* **18**, 559–563 (2017).
47. Lin, D. *et al.* Functional identification of an aggression locus in the mouse hypothalamus. *Nature* **470**, 221–226 (2011).
48. Reza, A. M. Realization of the Contrast Limited Adaptive Histogram Equalization (CLAHE) for Real-Time Image Enhancement. *The Journal of VLSI Signal Processing-Systems for Signal, Image, and Video Technology* **38**, 35–44 (2004).
49. Golden, S. A., Covington, H. E., Berton, O. & Russo, S. J. A standardized protocol for repeated social defeat stress in mice. *Nature Protocols* **6**, 1183–1191 (2011).
50. Golden, S. A. *et al.* Compulsive Addiction-like Aggressive Behavior in Mice. *Biological Psychiatry* **82**, 239–248 (2017).
51. Berger, A. L. *et al.* The Lateral Habenula Directs Coping Styles Under Conditions of Stress via Recruitment of the Endocannabinoid System. *Biological Psychiatry* **84**, 611–623 (2018).

52. He, K., Gkioxari, G., Dollár, P. & Girshick, R. Mask R-CNN. *arXiv:1703.06870 [cs]* (2018).
53. Redmon, J. & Farhadi, A. YOLOv3: an incremental improvement. *arXiv:1804.02767 [cs]* (2018).
54. Dollár, P. *Piotr's Computer Vision Matlab Toolbox*.
55. Breiman, L. Random forests. *Machine learning* **45**, 5–32 (2001).
56. Probst, P. & Boulesteix, A.-L. To Tune or Not to Tune the Number of Trees in Random Forest. *Journal of Machine Learning Research* 1–18.
57. Chawla, N. V., Bowyer, K. W., Hall, L. O. & Kegelmeyer, W. P. SMOTE: Synthetic minority over-sampling technique. *JAIR* **16**, 321–357 (2002).
58. Batista, G. E. A. P. A., Prati, R. C. & Monard, M. C. A study of the behavior of several methods for balancing machine learning training data. *SIGKDD Explor. Newsl.* **6**, 20 (2004).
59. Saito, T. & Rehmsmeier, M. The Precision-Recall Plot Is More Informative than the ROC Plot When Evaluating Binary Classifiers on Imbalanced Datasets. *PLoS ONE* **10**, e0118432 (2015).
60. Powers, D. Evaluation: from Precision, Recall and F-measure to ROC, Informedness, Markedness and Correlation. *Journal of Machine Learning Technologies* **2**, 37–63 (2011).
61. Dolensek, N., Gehrlach, D. A., Klein, A. S. & Gogolla, N. Facial expressions of emotion states and their neuronal correlates in mice. *Science* **368**, 89–94 (2020).
62. Grant, E. & Mackintosh, J. A comparison of the social postures of some common laboratory. *Behaviour* **21**, 246–259 (1963).
63. Miczek, K. A., Maxson, S. C., Fish, E. W. & Faccidomo, S. Aggressive behavioral phenotypes in mice. *Behavioural Brain Research* **125**, 167–181 (2001).
64. Yang, Q. & Wu, X. 10 challenging problems in data mining research. *Int. J. Info. Tech. Dec. Mak.* **05**, 597–604 (2006).
65. Haibo He & Garcia, E. A. Learning from Imbalanced Data. *IEEE Trans. Knowl. Data Eng.* **21**, 1263–1284 (2009).
66. Galar, M., Fernandez, A., Barrenechea, E., Bustince, H. & Herrera, F. A Review on Ensembles for the Class Imbalance Problem: Bagging-, Boosting-, and Hybrid-Based Approaches. *IEEE Trans. Syst., Man, Cybern. C* **42**, 463–484 (2012).
67. López, V., Fernández, A., García, S., Palade, V. & Herrera, F. An insight into classification with imbalanced data: Empirical results and current trends on using data intrinsic characteristics. *Information Sciences* **250**, 113–141 (2013).
68. Krawczyk, B. Learning from imbalanced data: open challenges and future directions. *Prog Artif Intell* **5**, 221–232 (2016).
69. Arakawa, T. *et al.* A male-specific QTL for social interaction behavior in mice mapped with automated pattern detection by a hidden Markov model incorporated into newly developed freeware. *Journal of Neuroscience Methods* **234**, 127–134 (2014).
70. Graving, J. M. *et al.* DeepPoseKit, a software toolkit for fast and robust animal pose estimation using deep learning. *eLife* **8**:e47994 (2019).
71. Arac, A., Zhao, P., Dobkin, B. H., Carmichael, S. T. & Golshani, P. DeepBehavior: A Deep Learning Toolbox for Automated Analysis of Animal and Human Behavior Imaging Data. *Front. Syst. Neurosci.* **13**, 20 (2019).
72. Liu, F. T., Ting, K. M. & Zhou, Z.-H. Isolation forest. in *2008 Eighth IEEE International Conference on Data Mining* 413–422 (IEEE, 2008). doi:10.1109/ICDM.2008.17.
73. Cao, Z., Simon, T., Wei, S.-E. & Sheikh, Y. Realtime multi-person 2D pose estimation using part affinity fields. *arXiv:1611.08050 [cs]* (2017).
74. Azevedo, A. W., Gurung, P., Venkatasubramanian, L., Mann, R. & Tuthill, J. C. A size principle for leg motor control in *Drosophila*. *bioRxiv* 730218 (2019).
75. Romero-Ferrero, F., Bergomi, M. G., Hinz, R. C., Heras, F. J. H. & de Polavieja, G. G. idtracker.ai: tracking all individuals in small or large collectives of unmarked animals. *Nat Methods* **16**, 179–182 (2019).
76. Pérez-Escudero, A., Vicente-Page, J., Hinz, R. C., Arganda, S. & de Polavieja, G. G. idTracker: tracking individuals in a group by automatic identification of unmarked animals. *Nat Methods* **11**, 743–748 (2014).
77. Brox, T., Rosenhahn, B., Cremers, D. & Seidel, H.-P. High accuracy optical flow serves 3-D pose tracking: exploiting contour and flow based constraints. in *Computer Vision – ECCV 2006* (eds. Leonardis, A., Bischof, H. & Pinz, A.) vol. 3952 98–111 (Springer Berlin Heidelberg, 2006).
78. He, K., Gkioxari, G., Dollár, P. & Girshick, R. Mask R-CNN. *arXiv:1703.06870 [cs]* (2018).
79. Tomaschek, F., Hendrix, P. & Baayen, R. H. Strategies for addressing collinearity in multivariate linguistic data. *Journal of Phonetics* **71**, 249–267 (2018).

80. Barga, R., Fontama, V., Tok, W. H. & Cabrera-Cordon, L. *Predictive analytics with Microsoft Azure machine learning*. (Springer, 2015).
81. Golovin, D. *et al.* Google vizier: A service for black-box optimization. in *Proceedings of the 23rd ACM SIGKDD international conference on knowledge discovery and data mining* 1487–1495 (2017).
82. Liberty, E. *et al.* Elastic Machine Learning Algorithms in Amazon SageMaker. in *SIGMOD'20* (June 14-19).
83. Feurer, M. *et al.* Efficient and robust automated machine learning. in *Advances in neural information processing systems* 2962–2970 (2015).
84. Pedregosa, F. *et al.* Scikit-learn: Machine learning in Python. *Journal of machine learning research* **12**, 2825–2830 (2011).
85. Christ, M., Braun, N., Neuffer, J. & Kempa-Liehr, A. W. Time Series Feature Extraction on basis of Scalable Hypothesis tests (tsfresh – A Python package). *Neurocomputing* **307**, 72–77 (2018).
86. Guyon, I. & Elisseeff, A. An introduction to variable and feature selection. *Journal of machine learning research* **3**, 1157–1182 (2003).
87. Branson, K., Robie, A. A., Bender, J., Perona, P. & Dickinson, M. H. High-throughput ethomics in large groups of *Drosophila*. *Nat Methods* **6**, 451–457 (2009).
88. Dankert, H., Wang, L., Hoopfer, E. D., Anderson, D. J. & Perona, P. Automated monitoring and analysis of social behavior in *Drosophila*. *Nat Methods* **6**, 297–303 (2009).
89. Jhuang, H. *et al.* Automated home-cage behavioural phenotyping of mice. *Nat Commun* **1**, 1–10 (2010).
90. de Chaumont, F. *et al.* Computerized video analysis of social interactions in mice. *Nature Methods* **9**, 410–417 (2012).
91. Burgos-Artizzu, X. P., Dollar, P., Dayu Lin, Anderson, D. J. & Perona, P. Social behavior recognition in continuous video. in *2012 IEEE Conference on Computer Vision and Pattern Recognition* 1322–1329 (IEEE, 2012). doi:10.1109/CVPR.2012.6247817.
92. Matsumoto, J. *et al.* A 3D-Video-Based Computerized Analysis of Social and Sexual Interactions in Rats. *PLOS ONE* **8**, e78460 (2013).
93. Berman, G. J., Choi, D. M., Bialek, W. & Shaevitz, J. W. Mapping the stereotyped behaviour of freely moving fruit flies. *J. R. Soc. Interface* **11**, 20140672 (2014).
94. Hong, W. *et al.* Automated measurement of mouse social behaviors using depth sensing, video tracking, and machine learning. *Proc Natl Acad Sci USA* **112**, E5351–E5360 (2015).
95. Wiltchko, A. B. *et al.* Mapping Sub-Second Structure in Mouse Behavior. *Neuron* **88**, 1121–1135 (2015).
96. Unger, J. *et al.* An unsupervised learning approach for tracking mice in an enclosed area. *BMC Bioinformatics* **18**, (2017).
97. Arakawa, T. *et al.* Automated Estimation of Mouse Social Behaviors Based on a Hidden Markov Model. in *Hidden Markov Models: Methods and Protocols* (eds. Westhead, D. R. & Vijayabaskar, M. S.) 185–197 (Springer New York, 2017). doi:10.1007/978-1-4939-6753-7_14.
98. Štíh, V., Petrucco, L., Kist, A. M. & Portugues, R. Stytra: An open-source, integrated system for stimulation, tracking and closed-loop behavioral experiments. *PLoS Comput Biol* **15**, e1006699 (2019).
99. Rodriguez, A. *et al.* ToxTrac: A fast and robust software for tracking organisms. *Methods in Ecology and Evolution* **9**, 460–464 (2018).
100. Rodriguez, A., Zhang, H., Klaminder, J., Brodin, T. & Andersson, M. ToxId: an efficient algorithm to solve occlusions when tracking multiple animals. *Sci Rep* **7**, 14774 (2017).
101. Thanos, P. K., Restif, C., O'Rourke, J. R., Lam, C. Y. & Metaxas, D. Mouse Social Interaction Test (MoST): a quantitative computer automated analysis of behavior. *J Neural Transm* **124**, 3–11 (2017).
102. Machado, A. S., Darmohray, D. M., Fayad, J., Marques, H. G. & Carey, M. R. A quantitative framework for whole-body coordination reveals specific deficits in freely walking ataxic mice. *eLife* **4**, e07892 (2015).
103. Günel, S. *et al.* DeepFly3D: A deep learning-based approach for 3D limb and appendage tracking in tethered, adult *Drosophila*. *bioRxiv* 640375 (2019) doi:10.1101/640375.
104. Gomez-Marin, A., Partoune, N., Stephens, G. J. & Louis, M. Automated Tracking of Animal Posture and Movement during Exploration and Sensory Orientation Behaviors. *PLOS ONE* **7**, e41642 (2012).
105. Ohayon, S., Avni, O., Taylor, A. L., Perona, P. & Roian Egnor, S. E. Automated multi-day tracking of marked mice for the analysis of social behaviour. *Journal of Neuroscience Methods* **219**, 10–19 (2013).
106. Tungtur, S. K., Nishimune, N., Radel, J. & Nishimune, H. Mouse Behavior Tracker: An economical method for tracking behavior in home cages. *BioTechniques* **63**, 215–220 (2017).