

# 1   **SegColR: Deep Learning for Automated Segmentation and Color Extraction**

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## 6   **Abstract**

7   Citizen science platforms like iNaturalist generate biodiversity data at an unprecedented scale, with  
8   observations on the order of hundreds of millions. However, extracting phenotypic information from  
9   these images, such as color of organisms, at such a large scale poses unique challenges for biologists.  
10   Some of the challenges are that manual extraction of phenotypic information can be subjective and  
11   time-consuming. Fortunately, with the maturation of computer vision and deep learning, there is an  
12   opportunity to automate large parts of the image processing pipeline. Here, I present SegColR, a user-  
13   friendly software package that leverages two state-of-the-art deep learning models - GroundingDINO  
14   and SegmentAnything - to enable automated segmentation and color extraction from images. The  
15   SegColR package provides an R-based interface, making it more accessible to evolutionary biologists  
16   and ecologists who may not have extensive coding experience. The SegColR pipeline allows users to  
17   load images, automatically segment them based on text prompts, and extract color information from the  
18   segmented regions. The package also includes visualization and data summarization functions to  
19   facilitate downstream analysis and interpretation of the results.

## 20   **Key Words**

21   color, color pattern, citizen science, image segmentation, deep learning, iNaturalist

## 22 INTRODUCTION

23 Color is an important trait for many organisms, influencing various aspects of their ecology and  
24 physiology. It plays a crucial role across the tree of life in communication, pollination syndromes,  
25 mating behavior, predator-prey interactions, and thermal regulation (e.g., Endler 1993; Huyghe et al.  
26 2007; Tsuchida et al. 2010; Finkbeiner et al. 2014; Mitchem 2017; Narbona et al. 2021; Cox and Davis  
27 Rabosky 2023). However, obtaining large-scale, high-quality color data can be a significant challenge  
28 for researchers. For example, while research museum collections house vast amounts of biodiversity  
29 information, the preservation techniques used may distort the original coloration of specimens (Pohland  
30 and Mullen 2006). Nonetheless, the ubiquity and importance of color in organismal biology means that  
31 accurately quantifying color traits is an important step towards understanding many aspects of ecology  
32 and evolution of organisms.

33 Citizen science initiatives like iNaturalist generate data in the form of crowdsourced  
34 observations and images, centralizing vast quantities of biodiversity data and providing opportunity to  
35 study color variation across the tree of life and at large spatial scales. Yet, the sheer scale of the data  
36 generated and the fact that it is typically not standardized, make it difficult to automate data collection  
37 and extraction. As such, most attempts to utilize resources like iNaturalist rely on manually extracting  
38 information image by image. Though this approach almost certainly results in high-quality datasets, for  
39 even relatively simple tasks like color quantification, extracting information from images will be time-  
40 consuming and likely require large teams of willing annotators. Furthermore, manual color extraction  
41 can be subject to measurement error, e.g. what one observer calls white another may call light green.  
42 Fortunately, the maturation of computer vision and deep learning techniques offers a promising  
43 solution to this challenge. By leveraging state-of-the-art machine learning models, it is now possible to  
44 automate large parts of the image processing pipeline, allowing biologists to extract phenotypic  
45 information from citizen science data at unprecedented scales.

46 Many of the most successful deep learning applications in recent years have been for computer  
47 vision tasks (LeCun et al. 2015; Goodfellow et al. 2016). Two key areas of computer vision that are  
48 particularly relevant for extracting color data from biodiversity images are object detection and  
49 instance segmentation. Object detection algorithms aim to locate and classify distinct objects within an  
50 image (Redmon et al. 2016; Ren et al. 2017), while instance segmentation models, delineate pixel-level  
51 boundaries of each instance of an object (He et al. 2017; Kirillov et al. 2020). By combining these  
52 capabilities, it becomes possible to focus color analysis only on the specific organisms of interest,  
53 rather than the entire scene. However, training a specialized computer vision model from scratch would

54 require large, annotated datasets. This would be prohibitively time-consuming to collect, especially for  
55 diverse taxonomic groups. Fortunately, progress in computer vision has led to the development of  
56 powerful pre-trained models that can be effectively leveraged for a variety of tasks with minimal fine-  
57 tuning (Zhai et al. 2022).

58 In this work, I utilize two such pre-trained models: GroundingDINO (Liu et al. 2024) for object  
59 detection and a data efficient version of the Segment Anything Model (SAM) (Kirillov et al. 2022;  
60 Chen et al. 2024) for instance segmentation. To provide an accessible entry point for ecologists and  
61 evolutionary biologists, I have implemented this deep learning-powered pipeline as both a set of  
62 Python scripts and an R package called SegColR. The R framework gives users who are already  
63 familiar with the R programming language an easier integration with their existing workflows and  
64 analytical tools. Additionally, the Python scripts are available for those researchers who may prefer a  
65 more customizable, low-level approach to image processing and analysis. The base SegColR package  
66 performs object detection and instance segmentation allowing for individual instances of focal taxa to  
67 be extracted from an image. These individual instances then allow for the extraction of color pixel-by-  
68 pixel. I demonstrate the SegColR workflow for several examples, showing also how one would assess  
69 the quality of the segmentation and color extraction. Finally, I outline potential pitfalls researchers may  
70 experience when using this and other automatic object detection and segmentation software.

## 71 **METHODS AND RESULTS**

### 72 **Object detection and instance segmentation**

73 Object detection and instance segmentation are fundamental tasks in computer vision (Chollet 2021).  
74 The goal of object detection is to draw bounding boxes around particular objects of interest and  
75 associate the bounding box with a particular class for a given image. However, most object detection  
76 algorithms are limited to a pre-determined set of classes. This is problematic for biological datasets as  
77 existing pre-trained models are unlikely to have been trained on all taxonomic groups of interest and  
78 adding new classes would require collecting and labeling new data in order to retrain the model.  
79 Attempts to address this challenge have focused on combining visual and textual modalities, with  
80 Grounding DINO successfully generalizing object detection (Liu et al. 2024). Grounding DINO is a  
81 transformed-based architecture that fuses language and vision modalities by linking the closed-set  
82 detector, DINO (Zhang et al. 2022), with grounded language-based pre-training (e.g., GLIP; Li et al.  
83 2022). The effect of this design is that Grounding DINO is able to detect arbitrary objects based on  
84 diverse text prompts. This model has been extended in several ways, including combining it with the

85 instance segmentation model, SegmentAnything (SAM; Kirillov et al. 2022). SAM, is an image  
86 segmentation model trained on the largest segmentation dataset with over 1 billion masks and 11  
87 million images. This allows it to achieve consistently high performance on zero-shot segmentation  
88 tasks even when compared to fully supervised models (Kirillov et al. 2022). The combination of  
89 Grounding DINO and SAM is called GroundedSAM (Ren. et al 2024) and it uses the bounding box  
90 output of Grounding DINO as the input of SAM for high quality instance segmentation. This approach  
91 can be further refined by using recently developed efficient versions of SAM such as SlimSAM, which  
92 achieve high accuracy while using far less training data (Chen et al. 2024). SlimSAM result in a model  
93 a fraction of the size of the original SAM (1.4% of the original parameters) and is ideally suited for  
94 biological research as the workflow can be run on moderately powerful personal computers.

## 95 **Color extraction**

96 The process of object detection and instance segmentation results in a set of masks for each instance of  
97 a particular class. Masks are logical matrices that represent the pixels within an image that correspond  
98 to a detected object. Each mask also has a confidence score indicating the model's certainty about the  
99 presence and classification of the detected object. When multiple detections are present for a single  
100 class, SegColR combines the masks that meet or exceed a user-specified score threshold. This is a  
101 quality control step that users are free to adjust based on the particular needs of their project. On one  
102 hand, higher confidence thresholds will be more conservative, but result in the inclusion of only the  
103 most reliable object detections. On the other hand, low confidence thresholds will include more  
104 detections, but there is a greater chance of inaccurate object detections.

105         Once the final mask for an object is obtained, SegColR extracts basic color information from  
106 the pixels in the original image. This is accomplished by converting the pixel values from the RGB  
107 color space to the Lab color space (CIE, 1976) and by default applying a k-means clustering algorithm  
108 (MacQueen 1967) to identify the dominant colors within the region of interest. If desired, it is possible  
109 to avoid k-means clustering by specifying a set of dominant colors which are then used to cluster each  
110 pixel based on the minimum euclidean distance between the pixel color and dominant colors within the  
111 Lab color space. The resulting dominant colors are characterized by their Lab coordinates and  
112 hexadecimal codes. Additional summary statistics, such as the mean and median color, are also  
113 calculated by default. More detailed color analysis is left to the user with other R packages providing  
114 more in-depth tool-kits once segmentation has been completed (e.g., Van Belleghem et al., 2018; Maia  
115 et al., 2019; van den Berg et al., 2020; Weller et al. 2024).

## 116 **SegColR features and examples**

### 117 *Description of the SegColR pipeline*

118 Using SegColR requires a user to input an image (specified by the path to the image) and a set of  
119 labels. The primary function of SegColR, `grounded_segmentation_cli` (Table 1), then performs  
120 grounded segmentation based on the input image and labels. This function creates a custom command-  
121 line interface (CLI) tool which interacts with a Python back-end. The Python back-end then uses the  
122 transformers library, which is itself an API to download and train pre-trained models, to call the  
123 segmenter and detector models (Wolf et al. 2020). Note that once the desired models are downloaded  
124 and necessary libraries installed, the SegColR software can be used entirely offline. Results are saved  
125 as a JSON file which can then be read into R using the function `load_segmentation_results`. Once the  
126 grounded segmentation is complete and the results are loaded into R, the segmentation can be plotted  
127 using `plot_seg_results` and a preliminary color analysis can be conducted using  
128 `process_masks_and_extract_colors` (Table 1). Finally, `plot_color_info` can be used to display several  
129 color summaries of the segmented image including dominant colors, mean and median colors, and  
130 RGB histograms. This pipeline is designed in the hope that it will be easy to execute and evaluate the  
131 resulting segmentation.

132         It is worth noting that a pre-requisite to using SegColR is having all the dependent Python  
133 libraries installed for a specified conda environment (`conda_env` argument of  
134 `grounded_segmentation_cli`). To assist users, I have created the function `setup_conda_environment`,  
135 which will install all the necessary libraries with either exact version numbers known to work with  
136 SegColR (`env_type= "specific"`) or with the minimum constraints on the required libraries (`env_type=`  
137 `"general"`). I recommend using `env_type= "general"` in most cases, noting that it takes longer to setup  
138 the environment than `env_type= "specific"` because library versions must be solved to ensure  
139 compatibility.

140 Table 1: Primary analysis functions.

Function	Argument	Definition
grounded_segmentation_cli	image_path	Character string. Path to the input image. E.g., “/home/user_name/project_directory/flower.jpg”
	labels	Character vector. Labels to detect in the image. E.g., c(“a flower”, “a tree”)
	threshold	Numeric. The minimum detection threshold for which detected objects will be segmented (default: 0.3).
	detector_id	Character string. ID of the detector model. default: “IDEA-Research/grounding-dino-tiny”
	segmenter_id	Character string. ID of the detector model. default: “Zigeng/SlimSAM-uniform-77”
	output_plot	Character string. Path or directory to save the output plot.
	output_json	Character string. Path or directory to save the output JSON.
	conda_env	Character string. Name of the conda environment to use. default: “segcolr-env” created by setup_conda_environment
load_segmentation_results	image_path	Character string. Path to the original image file.
	json_path	Character string. Path to the JSON file containing segmentation results. The JSON file is created by grounded_segmentation_cli
process_masks_and_extract_colors	image	Array. The original image (height x width x channels x 1).
	masks	List of logical matrices representing individual masks.
	scores	Numeric vector of scores corresponding to each mask.
	labels	Character vector of labels corresponding to each mask.
	include_labels	Character vector of labels to include.
	exclude_labels	Character vector of labels to exclude.
	score_threshold	Numeric. The score threshold for including a mask (default: 0.5).
	n_colors	Integer. Number of dominant colors to extract (default: 5).

### 141 Example 1 – Grounded segmentation and color analysis

142 This example demonstrates the utility of grounded segmentation in isolating organisms of interest from  
 143 background information. The subject of this analysis is an Andaman Hind fish, sourced from an  
 144 iNaturalist observation. Traditional color extraction methods applied to the entire image would yield  
 145 unusable data due to the background being present in the image. However, by employing grounded  
 146 segmentation, we can automatically focus on the organism of interest as informed by the user imputed  
 147 label, significantly improving the accuracy of color extraction. The grounded\_segmentation\_cli  
 148 function is utilized to segment the image, with parameters specifying the image path, label, and output  
 149 paths for JSON data and preliminary plot generation (Table 1).

```

150 ground_results <- grounded_segmentation_cli(
151   image_path = "/home/user/images/AndamanHind.jpeg",
152   labels = "a fish.",
153   output_json = "/home/user/output_dir/json/",
154   output_plot = "/home/user/SegColR/extdata/plot/")

```

155 The output JSON contains information on the score and label for all detected objects as well as the  
156 mask for that particular object. The segmentation results are then loaded and plotted using  
157 `load_segmentation_results` and `plot_seg_results` respectively (Figure 1). Following segmentation, the  
158 `process_masks_and_extract_colors` function is employed to extract color information from the focal  
159 organism. This function requires five key arguments, all of which are stored in the output of  
160 `grounded_segmentation_cli`. The results of this color analysis are then visualized using the  
161 `plot_color_info` function, which offers the option to recolor the mask based on the dominant colors  
162 extracted (Figure 2). Within R and for ease of use in downstream analysis, both the segmentation  
163 results and the color information results are list objects which containing information such as vectors of  
164 labels, confidence scores, masks, dominant colors, pixel by pixel coloration, mean and median colors.

```
165 color_results_k <- process_masks_and_extract_colors(  
166   image = ground_results$image,  
167   masks = ground_results$mask,  
168   scores = ground_results$score,  
169   labels = ground_results$label,  
170   include_labels = ground_results$label)
```

#### 171 *Example 2 – Multiple instances and excluding labels*

172 This example illustrates the capability of the segmentation algorithm to handle images containing  
173 multiple instances of a particular label and to exclude overlapping objects and organisms that may  
174 interfere with the color analysis. The focus here is on segmenting flowers to extract petal color, in an  
175 image that contains multiple flowers and a bee (Figure 3a). The segmentation process assigns scores to  
176 each detected instance, allowing for the exclusion of instances below a specified threshold. Setting this  
177 threshold to 0.5 results in the retention of two flower instances and one bee instance in the segmented  
178 result (Figure 3b). Other than using thresholds, a key feature of the `process_masks_and_extract_colors`  
179 function is its ability to exclude particular labels directly. In cases where multiple organisms are present  
180 in a scene, but color is desired from only one of them, specifying each object individually and then  
181 focusing the color extraction should produce higher quality results. This functionality is particularly  
182 valuable when combined with grounding DINO, as it provides remarkable flexibility in color extraction  
183 from non-standard images. In this case, it allows for the removal of the bee instance from the color  
184 extraction process (Figure 3c), ensuring that only the colors of the flowers are analyzed.

```
185 color_results <- process_masks_and_extract_colors(  
186   image = seg_results$image,  
187   masks = seg_results$mask,
```

```
188     scores = seg_results$score,  
189     labels = c("a flower.", "a bee."),  
190     include_labels = c("a flower."),  
191     exclude_labels = c("a bee."),  
192     score_threshold = 0.5,  
193     n_colors = 5  
194 )
```

### 195 *Example 3 – Specifying particular parts of an organism*

196 The final example demonstrates the power of grounding DINO in specifying and isolating particular  
197 parts of an organism for focused color analysis. This capability is important when color patterning is  
198 localized to specific areas of an organism. The subject of this analysis is a horned bream, where the  
199 main color patterning of interest is located on the body. The challenge lies in excluding the fins, which  
200 do not contain the coloration of interest. By leveraging the broad textual understanding provided by  
201 grounding DINO, we can use text prompts to detect and exclude various fin types from the analysis  
202 (Figure 4). This example highlights both the strengths and limitations of the current implementation.  
203 While the algorithm successfully identified and excluded the more prominent caudal and pelvic fins, it  
204 struggled with the detection of dorsal, anal, and pectoral fins. This suggests that, despite  
205 groundingDINO's generality, fine-tuning the model on domain-specific datasets will still be necessary  
206 in some cases.

```
207 ground_results <- grounded_segmentation_cli(  
208   image_path = "/home/user/project_file/img.jpg",  
209   labels = c("a fish.", "the fins of a fish."),  
210   output_json = "/home/user/project_file/json/",  
211   output_plot = "/home/user/project_file/plot/")
```

### 212 **Limitations**

213 The zero-shot object detection capabilities of groundingDINO (Feng et al., 2023) offer significant  
214 advantages when processing non-standardized citizen science data, particularly in the context of  
215 biodiversity studies. The heterogeneous nature of images on platforms like iNaturalist, where  
216 organisms may be positioned anywhere within the frame, camouflaged against diverse backgrounds, or  
217 present alongside other species, highlights the value of general text-prompt-based object detection. This  
218 approach greatly enhances the accessibility and usability of extensive biodiversity datasets compiled  
219 through citizen science initiatives. However, it is crucial to acknowledge that improved object detection  
220 algorithms cannot address all inherent biases present in citizen science data. A primary concern is the



221 inconsistency in lighting conditions across images, which can lead to shadow-induced distortions in  
222 color values (Szeliski, 2022). While some shadow removal techniques based on luminescence values  
223 have been developed (e.g., Murali & Govindan, 2013), their efficacy when applied to the diverse  
224 lighting conditions encountered in iNaturalist images has been limited. One potential approach to  
225 mitigate these lighting-related issues involves the use of pre-specified color palettes. Within the  
226 SegColR framework, this can be implemented through the `custom_colors` argument in the  
227 `process_masks_and_extract_colors` function. This method clusters pixel colors based on their distance  
228 to predefined custom colors rather than employing k-means clustering, potentially offering more  
229 consistent results across varying lighting conditions.

230         Beyond these technical limitations, citizen science datasets are subject to broader constraints  
231 that merit consideration. A significant issue is the presence of sampling effort biases (Dickinson et al.,  
232 2010). The majority of data is generated from North America, particularly the United States, despite the  
233 fact that global biodiversity is concentrated in tropical regions, which remain critically undersampled  
234 (e.g. Vasconcelos, 2023). While initiatives to increase sampling efforts in tropical areas exist, current  
235 datasets exhibit substantial geographical biases that must be accounted for in analyses utilizing citizen  
236 science data (Ward, 2014). Another limitation specific to color analysis is the discrepancy between  
237 human color perception (and standard camera capabilities) and the color perception of various  
238 organisms (Endler & Mielke, 2005). Many species perceive colors in ways that differ significantly  
239 from human vision, often requiring specialized equipment to capture color accurately. The widespread  
240 adoption of such specialized imaging technology among citizen scientists contributing to platforms like  
241 iNaturalist is unlikely, potentially limiting analyses to the human-perceptible color space. However, it is  
242 worth noting that images captured with specialized equipment could still be processed and analyzed  
243 using tools like SegColR, should they become available.

## 244 **DISCUSSION**

245 The proliferation of citizen science initiatives has led to an unprecedented accumulation of biodiversity  
246 data, offering researchers a vast repository of information. However, the heterogeneous nature of these  
247 datasets presents significant challenges for standardization and analysis. Recent advancements in  
248 computer vision technologies have emerged as a promising solution to address these complexities,  
249 offering remarkable flexibility in data processing and quantification. The application of deep-learning  
250 to tasks within ecology and evolutionary biology is not new, however, to-date, most applications have  
251 focused on models trained from scratch (e.g., Weaver and Smith 2023). A more powerful, accessible,

252 and environmentally responsible way forward for academics may be the utilization of large pre-trained  
253 models. These models are trained on extensive datasets which few academic collaborations are able to  
254 match (e.g., the billion masks used to train SAM; Krillov et al., 2023). Fortunately, many of the  
255 underlying representations within the hidden layers of the deep-learning models are still useful in  
256 biological studies and through the use of fine-tuning, can be readily adapted to even the most obscure  
257 taxonomic groups.

258         Several R-packages exist for color and color pattern analysis (e.g., Van Belleghem et al., 2018;  
259 Maia et al., 2019; van den Berg et al., 2020; Weller et al. 2024). These packages have several novel  
260 analysis features and users are encouraged to use SegColR in conjunction with existing R-packages  
261 rather than the final end-point of data collection. Color extraction from natural images poses several  
262 challenges including differentiating the signal of the focal organism from the natural background (see  
263 2.4 Limitations). SegColR uses object detection via groundingDINO to overcome this, but other  
264 approaches should be used in conjunction to ensure a high quality dataset (e.g., van den Berg et al.,  
265 2020). Other issues, such as the shadow distorting color, are more difficult to resolve. Using a pre-  
266 defined color palette with colors that are distant in color space may be one way around this, but shadow  
267 removal techniques are also a promising way forward (e.g., Murali & Govindan, 2013).

268         The implementation of deep learning tools in R is an important step towards making these  
269 techniques widely available to biologists. R is one of the most widely used programming language for  
270 academic ecologists and evolutionary biologists, but most deep-learning developments take place in  
271 Python. Furthermore, the use of light-weight models, such as slimSAM (Chen 2024), allows for  
272 advanced deep-learning models to be run even on moderately powerful personal computers. By  
273 enhancing the accessibility of these tools, biologists can gain access to an increasing number of data  
274 sources. Nonetheless the automation provided by computer vision techniques will need to be balanced  
275 with careful verification of the results. Performance is not guaranteed to be the same across all images  
276 (see example 2.3.3) and increasing the accuracy of these models on diverse taxonomic groups will  
277 likely require some amount of additional data collection and fine-tuning. Furthermore, while SegColR  
278 allows for a more automated collection of citizen science data, it cannot address all inherent limitations  
279 of this data source. Researchers must remain cognizant of lighting inconsistencies and sampling biases  
280 when interpreting results derived from these datasets.

## 281 CONCLUSION

282 Color is a crucial phenotype for many organisms, however gathering large interspecific datasets has  
283 proven difficult without some amount of automation. Here I have introduced SegColR, an R package  
284 which utilizes an underlying Python framework to automatically detect and segment organisms within  
285 citizen science photos. Using the pre-trained models groundingDINO and slimSAM, SegColR is able  
286 to preform object detection and segmentation without training for specific taxa. Using a computer  
287 vision pipeline for automated color extraction is just one example of how deep learning techniques can  
288 be used to quantify large-scale bio-diversity datasets. SegColR and other deep-learning software offer  
289 many exciting possibilities, but still necessitate careful consideration of the data produced. Researchers  
290 must reconcile the fact that though these tools are able to automatically extract vast quantities of data,  
291 high quality datasets will only be created if they are rigorously evaluated. As deep-learning techniques  
292 continue to integrate with ecology and evolutionary biology, it is important that they are used to  
293 complement traditional methods, rather than replace them.

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## 298 Data Availability Statement

299 The development version of the package is available on GitHub: <https://github.com/jboyko/SegColR>.  
300 All images and code used to generate the examples are taken from the vignette associated with the  
301 package. Data and image credits can be loaded using SegColR::load\_segcolr\_example\_data.

## 302 Conflict of Interest Statement

303 The author declares no competing interests.

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