

NEON USER GUIDE TO TICKS SAMPLED USING DRAG CLOTHS (NEON.DP1.10093)

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CHANGE RECORD

REVISION	DATE	DESCRIPTION OF CHANGE
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1 DESCRIPTION

1.1 Purpose

This document provides an overview of the data included in this NEON Level 1 data product, the quality controlled product generated from raw Level 0 data, and associated metadata. In the NEON data products framework, the raw data collected in the field, for example, the dry weights of litter functional groups from a single collection event are considered the lowest level (Level 0). Raw data that have been quality checked via the steps detailed herein, as well as simple metrics that emerge from the raw data are considered Level 1 data products.

The text herein provides a discussion of measurement theory and implementation, data product provenance, quality assurance and control methods used, and approximations and/or assumptions made during L1 data creation.

1.2 Scope

This document describes the steps needed to generate the L1 data product Ticks sampled using drag cloths (NEON.DP1.10093) - the abundance and density of ticks collected by drag and/or flag sampling, enumerated by species and/or lifestage - and associated metadata from input data. This document also provides details relevant to the publication of the data products via the NEON data portal, with additional detail available in the file, NEON Data Publication Workbook for Tick Sampling (NEON.DP1.10093) (AD[05]), provided in the download package for this data product.

This document describes the process for ingesting and performing automated quality assurance and control procedures on the data collected in the field pertaining to TOS Protocol and Procedure: Tick and Tick-Borne Pathogen Sampling (AD[07]). The raw data that are processed in this document are detailed in the file, NEON Data Ingest Workbook for Tick Sampling (NEON.DP0.10093) (AD[04]), provided in the download package for this data product. Please note that raw data products (denoted by 'DP0') may not always have the same numbers (e.g., '10033') as the corresponding L1 data product.

2 RELATED DOCUMENTS AND ACRONYMS

2.1 Associated Documents

AD[01]	NEON.DOC.000001	NEON Observatory Design (NOD) Requirements
AD[02]	NEON.DOC.000913	TOS Science Design for Spatial Sampling
AD[03]	NEON.DOC.002652	NEON Level 1, Level 2 and Level 3 Data Products Catalog
AD[04]	NEON.DP0.10093.001_dataValidation.csv	NEON Data Ingest Workbook for Tick Sampling (NEON.DP0.10093)
AD[05]	NEON.DP1.10093.001_variables.csv	NEON Data Publication Workbook for Tick Sampling (NEON.DP1.10093)
AD[06]	NEON.DOC.000911	TOS Science Design for Vectors and Pathogens
AD[07]	NEON.DOC.014045	TOS Protocol and Procedure: Tick and Tick-Borne Pathogen Sampling
AD[08]	NEON.DOC.000913	TOS Science Design for Spatial Sampling
AD[09]	NEON.DOC.000913	TOS Science Design for Plant Diversity
AD[10]	NEON.DOC.000008	NEON Acronym List
AD[11]	NEON.DOC.000243	NEON Glossary of Terms
AD[12]	OS_Generic_Transitions.pdf	NEON Algorithm Theoretical Basis Document: OS Generic Transitions
AD[13]		NEON's Ingest Conversion Language (NICL) specifications

3 DATA PRODUCT DESCRIPTION

The ticks sampled using drag cloths product provides counts of ticks collected using drag and/or flag processing. The drag method involves pulling a 1m² cloth along the ground at a slow pace (Milne 1943, Falco and Fish 1992). Where thick vegetation prevents continuous drag sampling, flag sampling is used instead. In flag sampling, the 1m² cloth is waved over and underneath vegetation rather than dragged along the ground (Ginsberg and Ewing 1989, Rulison et al. 2013). Collected ticks are enumerated by life-stage by NEON field technicians, after which tick samples are sent to one or more external facilities where a subset of collected ticks are identified to the lowest taxonomic rank possible (preferably species). Metrics of total tick abundance should be assessed from the field collection data, as counts of ticks by the taxonomist represent a subsample from the total number of ticks collected. From that subset of identified ticks, a portion are tested to quantify the prevalence of infection by various pathogens (see Tick-borne pathogen status NEON.DP1.10093.001). Some ticks are set aside for long-term archiving.

Ticks transmit numerous pathogens of wildlife, livestock, and humans, including the etiological agent of Lyme disease (*Borrelia burgdorferi*), the most frequently reported vector-borne disease of humans in the United States. Among arthropod vectors, ticks are particularly sensitive to meteorological conditions and associated physiological constraints, making it highly likely that the demography and biogeography of many tick species, and the pathogens they transmit, will be affected by climate change. Further, the multi-host lifecycles of most tick species increase their ecological connectivity and sensitivity to community-level perturbations that may arise from changes in human land- and resource-use practices.

3.1 Spatial Sampling Design

Tick sampling is conducted at regular intervals by NEON field technicians at terrestrial sites. At each site, six plots are selected for sampling, distributed within the sites proportional to permitted sampling area contained within each National Land Cover Database (NLCD) class. Ticks plots are situated such that the edge of each tick plot must be >150m from other NEON plots and infrastructure; tick plot centers are >500m from each other and no streams bisect tick plot perimeters. To reduce the probability that the sampling activities in other protocols impact tick diversity and distribution (e.g., technicians inadvertently attracting or redistributing ticks), tick plots centers are offset from distributed plots center according to a specified distance (150m +/- 15m) in a randomly chosen direction established prior to establishment of plots in the field. Sampling occurs by drag or flag sampling the 160m perimeter of each of the six 40m x 40m tick plots (Figure 1). See TOS Science Design for Vectors and Pathogens (AD[06]), TOS Protocol and Procedure: Tick and Tick-Borne Pathogen Sampling (AD[07]), TOS Science Design for Spatial Sampling (AD[08]) and TOS Science Design for Plant Diversity (AD[09]) for further details.

An exception to the design above occurs at NEON's Guanica Forest site (GUAN). At GUAN, the density of vegetation precluded sampling in any of the randomly selected plots using the above design. Instead, field staff located and marked six 'transect paths' where sampling is feasible. Each transect path contains between 80m² and 160m² of sampled area. Dragging and/or flagging then occurs over one more 10m long segments within each plot, with each segments generally within of another segment. Where possible, transect paths are located within 300m from the center of a mammal plot; 300m from the center of a bird plot; and 300m from the center of a Distributed Base Plot, with preference given to co-location with mammal plots where all three co-location criteria cannot be met.

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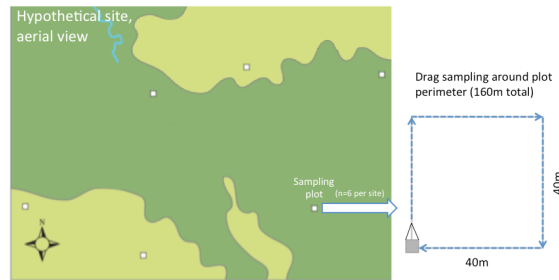


Figure 1: Drag and/or flag sampling occurs around the 160m perimeter of a 40mx40m plot

3.2 Temporal Sampling Design

Ticks are sampled during the growing season (determined by site-specific average greening period). During the growing season, sampling is conducted every three weeks at sites where ticks have previously been detected and every six weeks elsewhere. Sampling also occurs only if the high temperature on two consecutive days prior to planned sampling exceeds 0°C. For additional details, see the TOS Protocol and Procedure: Tick and Tick-Borne Pathogen Sampling.

3.3 Variables Reported

All variables reported from the field or laboratory technician (L0 data) are listed in the file, NEON Data Ingest Workbook for Tick Sampling (NEON.DP0.10093) (AD[04]). All variables reported in the published data (L1 data) are also provided separately in the file, NEON Data Publication Workbook for Tick Sampling (NEON.DP1.10093) (AD[05]).

Field names have been standardized with Darwin Core terms (<http://rs.tdwg.org/dwc/>; accessed 16 February 2014), the Global Biodiversity Information Facility vocabularies (<http://rs.gbif.org/vocabulary/gbif/>; accessed 16 February 2014), the VegCore data dictionary (<https://projects.nceas.ucsb.edu/nceas/projects/bien/wiki/VegCore>; accessed 16 February 2014), where applicable. NEON TOS spatial data employs the World Geodetic System 1984 (WGS84) for its fundamental reference datum and GEOID09 for its reference gravitational ellipsoid. Latitudes and longitudes are denoted in decimal notation to six decimal places, with longitudes indicated as negative west of the Greenwich meridian.

Some variables described in this document may be for NEON internal use only and will not appear in downloaded data.

3.4 Spatial Resolution and Extent

The finest resolution at which spatial data are reported is a single plot.

plotID (ID of plot within site) → **siteID** (ID of NEON site) → **domainID** (ID of a NEON domain).

The basic spatial data included in the data downloaded include the latitude, longitude, and elevation of the centroid of the plot where sampling occurred + associated uncertainty due to GPS error and plot width. Shape-

files of all NEON Terrestrial Observation System sampling locations can be found in the Document Library: <http://data.neonscience.org/documents>.

3.5 Temporal Resolution and Extent

The finest resolution at which temporal data are reported is the range between the startDate and endDate of each tick collection event (the duration during which dragging and/or flagging a single plot occurred).

The NEON Data Portal provides data in monthly files for query and download efficiency. Queries including any part of a month will return data from the entire month. Code to stack files across months is available here: <https://github.com/NEONScience/NEON-utilities>

3.6 Associated Data Streams

subsampleID is a linking variable that ties specific tick identifications and associated metadata to the Tick-borne pathogen status data product (DP1.10092). Visual assessments of the presence of ectoparasites, by life stage, on captured mammals can be found in the Small mammal box trapping data product (DP1.10072)

3.7 Product Instances

There are a maximum of 11 tick sampling bouts per year, with ticks collected from six plots per bout, yielding up to 66 tick samples per site per year. The total number of ticks per sampling event varies with site-specific conditions.

3.8 Data Relationships

The protocol dictates that each plot is sampled once per sampling bout (six expected records per eventID in tck_fielddata). A record from tck_fielddata may have zero (if no ticks present) or more child records in tck_taxonomyProcessed. An equivalent number of records will occur in tck_taxonomyRaw. A record from tck_taxonomyProcessed may have zero (if the pertinent subsampleIDs are not sent for pathogen testing) or more records in the tck_pathogen table in the Ticks and tick pathogens data product. Duplicates and/or missing data may exist where protocol and/or data entry aberrations have occurred; users should check data carefully for anomalies before joining tables.

tck_fielddata.csv (NEON.DP1.10093) - > One record expected per plotID for each eventID, generates a single sampleID if ticks are present

tck_taxonomyProcessed.csv (NEON.DP1.10093) - > Multiple record expected per sampleID, generates a unique subsampleID for each species*life stage combination found per sampleID. The taxonomic nomenclature in this file has been standardized and desynonymized according to NEON's master taxonomy for ticks.

tck_taxonomyRaw.csv (NEON.DP1.10093 - expanded package) - > Multiple record expected per sampleID, generates a unique subsampleID for each species*life stage combination found per sampleID. The taxonomic nomenclature in this file reflects the verbatim identifications provided by the external taxonomist and may contain synonyms.

tck_pathogen.csv (NEON.DP1.10092) - > Multiple record expected per subsampleID, generates a unique testingID for each tick selected for pathogen testing. Note not all subsampleIDs contribute to testing; some are simply archived in 95% EtOH.

tck_pathogenqa.csv (NEON.DP1.10092 - expanded) - > One record expected per batchID, contains relevant batch-level quality assurance procedures for each batch tested over all dates testingID for each tick selected for pathogen testing. Note not all subsampleIDs contribute to testing; some are simply archived in 95% EtOH.

4 DATA QUALITY

4.1 Data Entry Constraint and Validation

Many quality control measures are implemented at the point of data entry within a mobile data entry application or web user interface (UI). For example, data formats are constrained and data values controlled through the provision of dropdown options, which reduces the number of processing steps necessary to prepare the raw data for publication. An additional set of constraints are implemented during the process of ingest into the NEON database. The product-specific data constraint and validation requirements built into data entry applications and database ingest are described in the document NEON Data Ingest Workbook for Tick Sampling (NEON.DP0.10093), provided with every download of this data product. Contained within this file is a field named 'entryValidation-RulesForm', which describes syntactically the validation rules for each field built into the data entry application. Data entry constraints are described in NiCl syntax in the validation file provided with every data download, and the NiCl language is described in NEON's Ingest Conversion Language (NICL) specifications ([AD[13]).

Data collected prior to 2017 were processed using a paper-based workflow that did not implement the full suite of quality control features associated with the interactive digital workflow.

4.2 Automated Data Processing Steps

Following data entry into a mobile application or web user interface, the steps used to process the data through to publication on the NEON Data Portal are detailed in the NEON Algorithm Theoretical Basis Document: OS Generic Transitions (AD[12]).

4.3 Data Revision

All data are provisional until a numbered version is released; the first release of a static version of NEON data, annotated with a globally unique identifier, is planned to take place in 2020. During the provisional period, QA/QC is an active process, as opposed to a discrete activity performed once, and records are updated on a rolling basis as a result of scheduled tests or feedback from data users. The Change Log section of the data product readme, provided with every data download, contains a history of major known errors and revisions.

4.4 Quality Flagging

The **dataQF** field in each data record is a quality flag for known errors applying to the record. Please below for an explanation of **dataQF** codes specific to this product.

fieldName	value	definition
dataQF	legacyData	Data recorded using a paper-based workflow that did not implement the full suite of quality control features associated with the interactive digital workflow

5 REFERENCES

Falco, R. C., and D. Fish. 1992. A comparison of methods for sampling the deer tick, *Ixodes dammini*, in a Lyme disease endemic area. *Experimental & Applied Acarology* 14:165–173.

Ginsberg, H. S., and C. P. Ewing. 1989. Comparison of flagging, walking, trapping, and collecting from hosts as sampling methods for northern deer ticks, *Ixodes dammini*, and lone star ticks, *Amblyomma americanum* (Acari, Ixodidae). *Experimental and Applied Acarology* 7:313–322.

Milne, A. 1943. The comparison of sheep tick populations (*Ixodes ricinus* L). *Annals of Applied Biology* 30:240–250.

Rulison, E. L., I. Kuczaj, G. Pang, G. J. Hickling, J. I. Tsao, and H. S. Ginsberg. 2013. Flagging versus dragging as sampling methods for nymphal *Ixodes scapularis* (Acari: Ixodidae). *Journal of Vector Ecology* 38:163–167.