Empirical assessment of RAD sequencing for interspecific phylogeny
Astrid Cruaud, Mathieu Gautier, Maxime Galan, Julien Foucaud, Laure Sauné, Guénaëlle Genson, Emeric Dubois, Sabine Nidelet, Thierry Deuve, Jean Yves Rasplus

To cite this version:

HAL Id: hal-02638415
https://hal.inrae.fr/hal-02638415
Submitted on 11 Jun 2021

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L’archive ouverte pluridisciplinaire HAL, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d’enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

Distributed under a Creative Commons Attribution 4.0 International License
Empirical Assessment of RAD Sequencing for Interspecific Phylogeny

Astrid Cruaud,*†,1 Mathieu Gautier,‡,1,2 Maxime Galan,1 Julien Foucaud,1 Laure Sauné,1 Gwenâëlle Genson,1 Emeric Dubois,3 Sabine Nidelet,3 Thierry Deuve,4 and Jean-Yves Rasplus1
1INRA, UMR1062 CBGP, Montferrier-sur-Lez, France
2Institut de Biologie Computationnelle, 95 rue de la Galéria, 34095 Montpellier, France
3Montpellier GenomiX, c/o Institut de Génomique Fonctionnelle, Montpellier, France
4MNHN, UMR7205 OSEB, Muséum National d’Histoire Naturelle, Paris, France

*Corresponding author: E-mail: cruaud@supagro.inra.fr.
†These authors contributed equally to this work.

Associate editor: Emma Teeling

Abstract

Next-generation sequencing opened up new possibilities in phylogenetics; however, choosing an appropriate method of sample preparation remains challenging. Here, we demonstrate that restriction-site-associated DNA sequencing (RAD-seq) generates useful data for phylogenomics. Analysis of our RAD library using current bioinformatic and phylogenetic tools produced 400× more sites than our Sanger approach (2,262,825 nt/species), fully resolving relationships between 18 species of ground beetles (divergences up to 17 My). This suggests that RAD-seq is promising to infer phylogeny of eukaryotic species, though potential biases need to be evaluated and new methodologies developed to take full advantage of such data.

Key words: phylogenomics, next-generation sequencing, restriction-site associated DNA sequencing, empirical data, Insecta, Carabidae.

Next-generation sequencing opened up new possibilities for phylogenetics, allowing rapid and cost-effective generation of millions of reads from different loci on nonmodel species; however, choosing an appropriate method of library construction remains challenging. Consequently, Sanger sequencing of a few genes is still widely used to infer species phylogenies (McCormack et al. 2013). Sequencing complete eukaryotic genomes to infer species phylogenies remains expensive, time consuming, and unrealistic. Sequencing mitochondrial genomes is more affordable, but trees can be misleading due to introgression and heteroplasmy. Thus, methods based on reduced representations of the nuclear genome appear most adequate to generate large amounts of data across many individuals at reasonable costs.

Restriction-site-associated DNA sequencing (RAD-seq, Baird et al. 2008; supplementary fig. S1, Supplementary Material online) has been used to infer the recent evolutionary history (<3 My) of few organisms (e.g., Jones et al. 2013; Nadeau et al. 2013). However, with increasing genetic distances, mutations in the restriction sites may reduce the number of orthologous loci, making RAD-seq inappropriate to infer deeper relationships (McCormack et al. 2012). Conversely, recent in silico studies suggested that a sufficient number of markers could be obtained from distant species (up to 60 My old, Rubin et al. 2012). Here, we empirically tested this prediction by comparing the power of Sanger and RAD sequencing approaches to resolve relationships between 18 nonmodel species of ground beetles (Carabus), whose divergences ranged from 1.2 to 17 My (supplementary table S1, Supplementary Material online).

Details regarding data generation and analysis are described in the supplementary materials, Supplementary Material online.

Sanger—After a lengthy process of screening loci for variability, primer design and PCR optimization, maximum likelihood (ML) analyses of sequences from three mitochondrial and six nuclear markers led to poorly resolved and conflicting topologies, highlighting possible mitochondrial introgression between species (fig. 1A and B).

RAD-seq—After 4 days of library preparation following the protocol by Etter et al. (2011) (PstI enzyme), 2 weeks of sequencing on one lane of a HiSeq 2000 flowcell and a week of data processing on a standard computer (using Stacks; Catchen et al. 2011), we obtained 400× the volume of Sanger data and 270× more informative sites. Our data set resulted from a stringent loci selection to ensure for homology and minimize the amount of missing data. More than half of the individuals should have sequences for a loci to be included in our analysis, and the number of mismatches allowed when merging loci from all individuals varied from 4 to 10 (parameter n, cstacks).

Whatever the value of n, ML analysis of the RAD-seq data sets (up to 25,425 loci, i.e., one every 12,000 nt, supplementary table S2, Supplementary Material online) produced the same fully resolved topology (fig. 1C and supplementary fig. S2, Supplementary Material online).
different from the nuclear and mitochondrial Sanger trees. These two last topologies were rejected by statistical tests \( P < 0.001 \). Although significant loss of RAD markers occurred for the oldest DNA sample (78.3%–75.4%, Carabus olympiae, extraction performed in 1998), enough signal remained for its placement. Loss of RAD markers also occurred with increasing genetic distances, though enough information was retained to accurately resolve the relationships within Carabus (Deuve et al. 2012). When \( n \) was set to 10, more polymorphic loci were included and missing data reached 68.6% for a divergence of 17 My (fig. 1C and supplementary fig. S2, Supplementary Material online). Preliminary tests are thus recommended to optimize the amount of allowed mismatches within a RAD locus to avoid loci overkill while ensuring loci homology across individuals. Nevertheless, percentages of missing data were comparable between Sanger and RAD-seq data sets (fig. 1D, supplementary table S2, Supplementary Material online). Furthermore, missing data did not bias reconstruction, as topologies obtained from the complete matrices (loci shared by all 18 species) were similar to those inferred from the incomplete matrices (supplementary fig. S3, Supplementary Material online).

Overall, this study illustrates the power of RAD-seq to infer shallow relationships, and we believe that this approach may be generalized to many groups (~90% of the insect genomes range between 100 and 800 Mb, supplementary fig. S5, Supplementary Material online). Here, we used only one partition and relied on current evolutionary models to analyze RAD-seq data. Testing alternative partitioning schemes or developing more appropriate models to deal with the specificities of such data (e.g., modeling gain/loss of restriction sites) might be promising to improve inferences.

Finally, RAD-seq definitely opens new avenues for phylogeneticists but possibly also a Pandora’s box of analytical issues. These issues will need to be explored to avoid being
mislead by systematic (Lemmon EM and Lemmon AR 2013) or other biases (Arnold et al. 2013), which may differ depending on the level of genetic divergence among samples.

Supplementary Material

Supplementary figures S1–S5 and tables S1 and S2 are available at Molecular Biology and Evolution online (http://www.mbe.oxfordjournals.org.)

Acknowledgments

The authors thank the SeqGen plateform (CeMEB Labex, France) for facilitating access to the Covaris S220 instrument and the CBGP HPC computational platform on which analyses were performed. The authors also thank E. Artige (CBGP) for her help on the field, I. Meusnier (CBGP) for her assistance for lab work, P.A. Gagnaire for P1 adapters, K. Gharbi and C. Eland (GenePool, UK) for their assistance with transferring RAD-seq protocol to CBGP and two anonymous reviewers for their valuable comments on a previous version of this manuscript. This work was based upon financial support received from the division “Plant Health and the Environment” of the French National Institute for Agricultural Research (INRA) and from the network “Library of Life” funded by the National Center for Scientific Research (CNRS), the National Museum of Natural History (MNHN), the French National Agronomic Institute (INRA), and the French Alternative Energies and Atomic Energy Commission (CEA) (Genoscope). Sanger sequences were deposited in GenBank under accession numbers KJ158754-KJ158836. RAD-seq data sets can be downloaded from http://www1.montpellier.inra.fr/CBGP/NGS/Files/Datasets_Cruaud_et_al_2014_RAD_MBE.zip.

References


