Research Articles

New resources for the forensic genetics community available on the NIST STRBase website

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Abstract

For more than a decade, the U.S. National Institute of Standards and Technology (NIST) has maintained the short tandem repeat DNA Internet database (STRBase), which is located at http://www.cstl.nist.gov/biotech/strbase/. The purpose of STRBase has been and continues to be an attempt to bring together the abundant literature and information in the forensic genetics field in a cohesive fashion to make current and future work easier. New materials are regularly added to expand the valuable information contained on the STRBase website.

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1. Introduction

In late 1996 and early 1997, while a postdoctoral researcher working with Dennis Reeder at the National Institute of Standards and Technology (NIST), I compiled the initial information for and launched the short tandem repeat DNA Internet database (STRBase). Many of the core materials came from my Ph.D. dissertation on STR typing by capillary electrophoresis, which was written in the summer of 1995. A comprehensive examination of more than 500 articles from the scientific literature on STRs was conducted to organize useful information from the rapidly growing field of forensic genetics. The initial information contained in STRBase was described previously [1–3]. Over the past decade, aggressive collection of articles on STRs from forensic journals and other sources has resulted in over 2900 publications, which are listed in the STRBase reference section (http://www.cstl.nist.gov/biotech/strbase/str_ref.htm). This comprehensive collection of publications on the subject has led to several reviews on forensic DNA analysis [4,5].

2. Materials and methods

Information on the STRBase website is contained in hypertext markup language (HTML) files that were created primarily using Microsoft FrontPage Software (Richmond, WA). Over 1000 files now exist containing 10,000 plus printed pages of information that are connected with approximately 4600 hyperlinks. An additional 2300 hyperlinks connects various information on STRBase to other Internet websites including 342 direct links to various organizations, journals, academic and forensic institutes, commercial sites, genetic genealogy labs, parentage testing labs, and legal sites dealing with forensic DNA (see http://www.cstl.nist.gov/biotech/strbase/weblink.htm). Thus, STRBase is not a true searchable “database” but rather a collection of information with interconnected files. Since July 2006, resources that are added to STRBase are now tracked on a “recent updates” page: http://www.cstl.nist.gov/biotech/strbase/updates.htm.
3. Results and discussion

3.1. STR fact sheets, multiplexes, and variant alleles

STR fact sheets are the centerpiece of STRBase and list information regarding genomic location, GenBank accession, repeat structure, reported PCR primer sets, observed allele sizes and sequence structure, commercially available allelic ladders, common multiplexes, and mutation rates. A total of 53 STR fact sheets including all 18 core or common loci used in current commercial STR kits [6] are available at http://www.cstl.nist.gov/biotech/strbase/str_fact.htm. In recent months, new fact sheets for SE33, D2S1338, D19S433, D1S1656, and DYS635 have been created and information on other loci brought fully up-to-date to enable coverage of all commercial STR kits.

Multiplex assay and kits are visually summarized by locus-specific allele size ranges and dye colors with each locus name hyperlinked to the appropriate STR fact sheet. Labs worldwide continue to contribute to knowledge regarding rare alleles such that we now have catalogued 364 variant alleles and 153 tri-allelic patterns.

3.2. Training materials and NIST project team output

Numerous PowerPoint slides, NIST publications and presentations, software programs, and other useful information are available for download and use by the forensic genetics community. Materials from over a dozen recent workshops consisting of thousands of slides covering capillary electrophoresis, low-copy number DNA testing, mixture interpretation, qPCR DNA quantitation, Y-chromosome and mtDNA analysis, and validation are available for use at http://www.cstl.nist.gov/biotech/strbase/training.htm.

3.3. Information and tools to aid PCR primer set concordance studies

A new section of STRBase was recently created to track null alleles detected through DNA testing with different primer sets (http://www.cstl.nist.gov/biotech/strbase/NullAlleles.htm). Thus far, 38 entries covering 12 different STR loci are listed from various published studies including a recent concordance study of the new MiniFiler kit with other commercial STR kits [7].

A new software program running in Microsoft Excel and entitled STR_MatchSamples was created by David Duewer (NIST Chemical Sciences and Technology Laboratory) that enables concordance checking with multiple data sets. This program is listed in Table 1 along with others that can be downloaded from STRBase.

4. Conclusions

STRBase has been well received and widely used by the forensic genetics community and we continue to create additional resources for our website. In 2005, NIST adopted STRBase as an official Standard Reference Database (SRD) giving further credence to the value of the information contained in our website.

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Conflict of interest

None.

References


