Evaluation of a Cost Effective In-House Method for HIV-1 Drug Resistance Genotyping Using Plasma Samples

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Abstract

Objectives: Validation of a cost effective in-house method for HIV-1 drug resistance genotyping using plasma samples.

Design: The validation includes the establishment of analytical performance characteristics such as accuracy, reproducibility, precision and sensitivity.

Methods: The accuracy was assessed by comparing 26 paired Virological Quality Assessment (VQA) proficiency testing panel sequences generated by in-house and ViroSeq Genotyping System 2.0 (Celera Diagnostics, US) as a gold standard. The reproducibility and precision were carried out on five samples with five replicates representing multiple HIV-1 subtypes (A, B, C) and resistance patterns. The amplification sensitivity was evaluated on HIV-1 positive plasma samples (n = 88) with known viral loads ranges from 1000–1.8 million RNA copies/ml.

Results: Comparison of the nucleotide sequences generated by ViroSeq and in-house method showed 99.41 ± 0.46 and 99.68 ± 0.35% mean nucleotide and amino acid identity respectively. Out of 135 Stanford HIVdb listed HIV-1 drug resistance mutations, partial discordance was observed at 15 positions and complete discordance was absent. The reproducibility and precision study showed high nucleotide sequence identities i.e. 99.88±0.10 and 99.82±0.20 respectively. The in-house method showed 100% analytical sensitivity on the samples with HIV-1 viral load >1000 RNA copies/ml. The cost of running the in-house method is only 50% of that for ViroSeq method (1128 vs 3008), thus making it cost effective.

Conclusions: The validated cost effective in-house method may be used to collect surveillance data on the emergence and transmission of HIV-1 drug resistance in resource limited countries. Moreover, the wide applications of a cost effective and validated in-house method for HIV-1 drug resistance testing will facilitate the decision making for the appropriate management of HIV infected patients.

Introduction

Close to 60% of those who require anti-retroviral treatment (ART) are already under ART [1]. The long term success of this programme will depend on the continued virus suppression as a result of anti-retroviral treatment. However, in resource-limited countries, HIV drug resistance testing is not generally available or it is too costly to be used in the routine monitoring of patients receiving ART. Monitoring the trends of HIV drug resistance among treatment failures is crucial as HIV drug resistance is a major threat with ability to negate the benefits accrued by the free ART programme. Detection and monitoring of HIV drug resistance by molecular genotyping is pivotal to ensure ongoing regimen efficacy. Therefore, the World Health Organization (WHO) recommends population-based surveillance and monitoring of HIV drug resistance in resource-limited settings [2,3]. The reported pattern and rates of transmitted and acquired drug-resistant HIV variants will collectively form the regional and global recommendations on which ART is to be maintained or changed in the first and second-line antiretroviral regimens [4]. The ViroSeq Genotyping System 2.0 (Celera Diagnostics, US) and TruGene are the two FDA-approved commercially available methods for HIV drug resistance testing [5,6]. The majority of the HIV-1 infected patients cannot afford the commercial HIV-1 drug resistance testing when they experience ART failure because of its high cost. Thus, there is a need for the development of a cost effective and efficient in-house method for HIV-1 drug resistance testing for application in resource limited settings.

The development of an in-house method has been done independently in each lab, usually requiring investment of considerable effort to optimize the procedures used [7–11]. In larger studies and surveys, it is important to have confidence that the results generated from different participating labs are of high quality and comparable to each other. One solution to this
problem would be to recommend all labs to use the same method, but this is not practical, given that the local difference in reagent supply, HIV subtypes, personnel training and the requirement to change the established procedure may vary. An alternative approach is to use a validated method that will ensure quality results.

The validation of an in-house method is now a prerequisite for testing any sample for World Health Organization (WHO) recommended surveillance and studies [12]. Most of the laboratories have validated the in-house method by the minimal criteria strategy which includes the comparison of the sequence of large numbers of specimens (e.g. 50 to 200) obtained by an in-house method with that obtained by previously validated commercially available FDA approved HIV genotyping method as “Gold Standard” [7,9,10]. Whereas this minimal validation procedure can describe the accuracy of the method, further assessment of the other analytical characteristics such as reproducibility, precision and analytical sensitivity is also required to ensure the reliable results. The acceptability of the performance of a method is determined by the analytical performance characteristics such as accuracy, reproducibility, precision, amplification sensitivity and specificity.

In this study, previous in-house method [13] was modified and evaluated by the analytical performance characteristics such as accuracy, reproducibility, precision, analytical sensitivity.

Materials and Methods

Samples

The validation of an in-house method was done according to the WHO guidelines [12] including participation in Virological Quality Assessment (VQA) HIV Genotypic Drug Resistance proficiency testing panels (VQA contract # NO1-AI-50044). The accuracy of the in-house method was assessed by comparing 26 paired VQA HIV Genotypic Drug Resistance proficiency testing panel sequences. The nucleotide/amino acid sequence obtained by the ViroSeq method (Celera Diagnostics, US) was considered as the gold standard with which the nucleotide/amino acid sequences obtained by the in-house method were compared.

The samples used in the validation study had similar characteristics to the samples which are tested routinely (sample type: plasma, genetic subtype: HIV-1 subtype C, A and B, viral load range >1000 copies/ml, resistance pattern: reverse transcriptase and protease inhibitor mutations). The reproducibility and the precision were carried out on five samples with five replicates each. All these samples were previously tested for HIV-1 drug resistance genotyping using the ViroSeq method and aliquots were stored at −70±5°C.

Validation criteria

Accuracy was defined as detection of 99% of known HIV-1 drug resistance mutations when compared with the results obtained by the ViroSeq Genotyping System 2.0 (Celera Diagnostics, US). The reproducibility and precision were defined as ≥98% nucleotide identities in ≥90% of pairwise comparisons, with the mixtures being counted as partially discordant. The nucleotide/amino acid sequences were analyzed under three categories: [10] (i) Concordant (if both the ViroSeq and in-house method gave the same nucleotide/amino acid). (ii) Partially discordant (if nucleotide base/amino acid mixture by one method but not by the other) and (iii) Discordant (if the two methods detected different nucleotide bases/amino acids).

Analytical Sensitivity

Since HIV-1 subtype C is the predominant subtype in India, the analytical sensitivity of the in-house method was tested by using two HIV-1 subtype C clinical samples with the viral load 75,520 copies/ml and 18,500 copies/ml determined by the COBAS Amplicor HIV-1 Monitoring Kit version 1.5 (Roche Diagnostics, Branchburg, New Jersey). The samples were serially diluted (the created copy number ranged from 4720 to less than 1000 HIV-1 RNA copies/ml) using HIV negative human plasma. Viral RNA was then extracted and the pol gene was amplified by the in-house method as described below.

The protocol was finalized before initiation of the evaluation of the in-house HIV drug resistance assay. As per the WHO guidelines, an acceptance criterion for each analytical performance characteristic was established in advance [12]. The protocol for the in-house method used for the validation is described below.

RNA Extraction

RNA extraction from plasma samples were performed using the NucliSSENS® easyMAG® (Biomerieux, Durham, NC) automated nucleic acid extraction system according to the manufacturer’s recommendations (NucliSSENS easyMAG user manual, v 1.1; BioMerieux, Boxtel, Netherlands). Five hundred microlitres of each sample was placed in the disposable sample vessel and the sample vessel was loaded onto the extractor. After the initial lysis incubation for 10 min, 50 μl of magnetic silica was added to each sample and the extractor was restarted. The samples were eluted in 50 μl of extraction buffer 3. All samples were transferred to a 1.5-ml microcentrifuge tube and stored at −70°C.

RT-PCR and Nested PCR

RT-PCR was performed as described previously [13]. The nested PCR was performed using the inner primer pair as mentioned in Table 1 to get an amplified fragment of 1204 bp. The amplification was done using Taq DNA polymerase (Genei, Bangalore India) 3 U/μl (1 μl) in a 10 × PCR buffer B (5 μl), 25 mM Mg2+ (4 μl), 2 mM dNTPs mix (Fermentas) (5 μl), 10 pmol of each primer and 6 μl of RT-PCR product. The final volume of 50 μl reaction was made up using DNase/RNase free water. The reaction conditions used for nested PCR were: initial denaturation at 94°C for 2 min followed by DNA amplification: 25 cycles at 94°C for 20 sec, 59°C for 45 sec, 72°C for 3 min and a final extension for 10 min at 72°C.

A GeneAmp PCR system 9700 thermal cycler (Applied Biosystem, CA, USA) was used for all PCRs. The results were checked by electrophoresis of the nested PCR products on 1% agarose (Genei, Bangalore, India) gels containing ethidium bromide (Sigma Aldrich, USA) and visualization of the amplified bands under UV light.

The amplified PCR products were purified using the QIAquick PCR purification kit (QiaGen Hilden, Germany) and eluted in 30 μl elution buffer. The purified PCR product was directly sequenced in both directions using BigDye Terminator Cycle Sequencing Ready Reaction kit version 3.1 (Applied Biosystem, CA, USA).

Sequence analysis

DNA sequencing was performed on 3100 DNA genetic analyzer (Applied Biosystem, CA, USA) with the six specific primers mentioned in Table 1. These sequencing primers provided overlapping, bidirectional sequences covering the whole protease region and partial reverse transcriptase region. The sequencing
reaction was performed in 10 μl volume containing 2 μl ready reaction mix, 4 pmol primers, 2 μl of 5× sequencing buffer, 2 μl of purified PCR product (30 ng/μl) and the volume was adjusted to 10 μl with DNase/RNase free water. The sequencing reaction was carried out using 25 linear amplification cycles of 96°C for 10 sec, 50°C for 5 sec, 60°C for 4 min. Unincorporated dNTPs were then removed by precipitation with 80% isopropanol and the dried pellet was resuspended in 10 μl of Hi-DiTM formamide (Applied Biosystem, CA, USA). The raw sequence data from ABI 3100 genetic analyzer was assembled, aligned and edited with the SeqScape v2.0 software (Applied Biosystem, CA, USA).

To rule out PCR contamination, the phylogenetic tree was generated by using the PhyML software to create a maximum likelihood tree [14]. The reference sequences were obtained from the Los Alamos HIV Database (www.hiv.lanl.gov). The sequence quality was also checked by Sequence Quality Assessment Tool (SQUAT) analysis software [15]. The final edited sequences were submitted to the “HIVdb Program: Sequence Analysis” in the Stanford University HIV drug resistance database for drug resistance interpretation [16]. Pairwise nucleotide sequence identity and discrepancy were analyzed using BioEdit v 7.0.9.0 [17]. The Rega HIV-1 subtyping tool Version 2.0 was used to determine the HIV-1 subtype [http://dbpartners.stanford.edu/RegaSubtyping/ Accessed on 14th May 2012].

Statistical analysis
Wilcoxon Signed-Rank test was used to analyze the difference in number of nucleotide mixtures detected between the ViroSeq and in-house method. The statistical significance was considered when P value was <0.05.

Results
Accuracy
All 26 paired VQA HIV Genotypic Drug Resistance proficiency testing panel samples were amplified and sequenced successfully using both the in-house and ViroSeq methods. The mean nucleotide identity was 99.41±0.46% (mean ± SD) among paired nucleotide sequences (Table 2). Wilcoxon signed-rank test was used to compare the in-house and ViroSeq method basecalling for mixed bases and revealed no significant difference (P, 0.382).

A total of 135 drug resistance-associated amino acid positions in protease (36 mutations) and reverse transcriptase (99 mutations) region were detected among 26 paired sequences. There was no complete nucleotide or amino acid discordance. The overall amino acid codon agreement was 99.68±0.35% among paired amino acid sequences.

The partial discordance due to synonymous/non-synonymous substitutions was observed at 15 amino acid positions including 2 positions in protease and 13 positions in the reverse transcriptase region (Table 3). Out of 15 partial discordant positions, in 13 positions mixtures of the wild type and the mutant virus codons were detected by the in-house method only, while the ViroSeq method detected either the wild type or mutated codon. In the remaining two positions, mixtures of the wild type and the mutant virus codons were detected by the ViroSeq method only, while the in-house method could detect only the wild type or mutated codon.

Reproducibility
All five replicates of five samples were amplified and sequenced successfully. The drug resistance mutation pattern and subtype distribution of samples selected for reproducibility study are given in Table 4. A total of 25 sequences were generated from five samples. The phylogenetic tree for all 25 sequences along with HIV-1 reference sequences (n = 56) was constructed by using the PhyML software to create a maximum likelihood tree [14]. The sequences confirmed the absence of cross-contamination or sample mix-up. The paired sequences obtained for each sample were more closely related to one another than to the sequences of any other sample (Fig. 1). Moreover, as expected, sequences from the same sample clustered together.

We found high nucleotide sequence identities ranging from 99.72% to 100.00% (99.85±0.10) (Table 4). The minor differences observed in sequence identity were caused by base mixtures (nucleotide mixture by one method but not by another). However, this difference did not translate into differences in amino acids.

A total of 35 drug resistance-related positions [PI mutations (7), NRTI mutations (16) and NNRTI mutations (12)] for each sequence and its replicates were also analyzed for reproducibility. All 35 drug resistance mutations were found to be reproducible with 100% concordance.

Precision
The precision study also showed high nucleotide sequence identity ranging from 98.49% to 100.00% (99.82±0.20) (Table 4).
NRTI mutations (12) and NNRTI mutations (6) were seen for five replicates of five samples analyzed for precision study. In addition, one sample showed an insertion at amino acid 35 in the protease for all replicates. All 34 drug resistance mutations and one insertion at amino acid 35 in the protease were found to be reproducible with 100% concordance. The drug resistance mutation pattern and subtype distributions of samples selected for precision study is given in Table 4.

**Amplification Sensitivity**

The amplification sensitivity of the in-house method was evaluated on HIV-1 positive plasma samples (n = 88) with known viral load ranging from 1000 to 1.8 million RNA copies/ml (see Table S1) for the viral copy number range, % amplification and subtype distribution). All the samples with viral load >1000 HIV-1 RNA copies/ml were amplified and sequenced successfully.

We have also evaluated the amplification sensitivity by serial dilution of two samples with high viral loads to achieve a range of copy numbers (4720, 2360, 1180, 590 RNA copies/ml) using plasma from HIV-1 negative donor followed by four replicate testings of each dilution. The amplification was seen in all four replicates of the sample with 1000 RNA copies/ml, whereas amplification was observed only in two out of four replicates with <1000 RNA copies/ml for both samples.

### Table 3. Drug resistance-associated amino acid positions in protease and reverse transcriptase from 26 proficiency testing panel plasma samples genotyped by the in-house and the ViroSeq methods.

| Amino Acid | Amino acid detected by the in-house method (# of samples) | Amino acid detected by the ViroSeq method (# of samples) | # of partially discordant mutations |
|------------|-----------------------------------------------------------|--------------------------------------------------------|-------------------------------------|
| Protease   |                                                           |                                                        |                                     |
| L10        | L (21), I (4), IL (1)                                      | L (21), I (5)                                          | 1                                   |
| L23        | L (23), I (3)                                             | L (23), I (3)                                          | 0                                   |
| L33        | L (23), F (3)                                             | L (23), F (3)                                          | 0                                   |
| M46        | M (23), L (3)                                             | M (23), L (3)                                          | 0                                   |
| L47        | I (25), IV (1)                                            | I (26)                                                 | 1                                   |
| L54        | I (23), V (3)                                             | I (23), V (3)                                          | 0                                   |
| A71        | A (21), V (2), T (1), TI (2)                              | A (21), V (2), T (1), TI (2)                           | 0                                   |
| G73        | G (24), GS (2)                                            | G (24), GS (2)                                         | 0                                   |
| V82        | V (23), A (3)                                            | V (23), A (3)                                          | 0                                   |
| N88        | N (23), G (3)                                            | N (23), G (3)                                          | 0                                   |
| L90        | L (21), M (5)                                            | L (21), M (5)                                          | 0                                   |
| Reverse Transcriptase |                                         |                                                        |                                     |
| A41        | M (19), L (7)                                            | M (19), L (7)                                          | 0                                   |
| A62        | A (21), V (3), AV (2)                                     | A (21), V (3), AV (2)                                  | 0                                   |
| K65        | K (24), KR (2)                                            | K (24), KR (2)                                         | 0                                   |
| D67        | D (18), N (5), DN (2)                                     | D (17), N (5), DN (2)                                  | 0                                   |
| T69        | T (24), IT (2)                                            | T (24), IT (2)                                         | 0                                   |
| K70        | K (24), R (2)                                            | K (24), R (2)                                          | 0                                   |
| L74        | L (21), V (2), I (1), IL (2)                              | L (23), V (1), I (1), IV (1)                           | 2                                   |
| V75        | V (22), T (2), AV (1), AITV (1)                           | V (22), T (2), AITV (2)                                 | 1                                   |
| V90        | V (23), I (2), IV (1)                                     | V (23), I (2), IV (2)                                  | 1                                   |
| L100       | L (23), I (3)                                            | L (23), I (3)                                          | 0                                   |
| K101       | K (23), Q (2), KPQT (1)                                   | K (24), Q (2)                                          | 1                                   |
| K103       | K (11), N (13), KR (2)                                    | K (13), N (13)                                         | 2                                   |
| V106       | V (22), M (4)                                            | V (22), M (4)                                          | 0                                   |
| E138       | E (24), A (2)                                            | E (24), A (2)                                          | 0                                   |
| V179       | V (24), D (2)                                            | V (24), D (2)                                          | 0                                   |
| M184       | M (10), V (8), L (3), MV (5)                              | M (14), V (8), L (3), MV(1)                            | 4                                   |
| Y188       | Y (24), C (1), CY (1)                                     | Y (24), C (2)                                          | 1                                   |
| L210       | L (23), W (3)                                            | L (23), W (3)                                          | 0                                   |
| T215       | T (19), Y (5), CS(1), ST (1)                              | T (20), Y (5), C (1)                                   | 1                                   |
| H221       | H (23), Y (3)                                            | H (23), Y (3)                                          | 0                                   |
| P225       | P (24), H (1), HP (1)                                     | P (24), H (1), HP (1)                                  | 0                                   |

Note: Partial discordant positions are shown in bold.
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| Sample ID | HIV-1 VL (Copies/ml) | HIV-1 Subtype | % Nucleotide sequence identity | Replicate Tests | No. of drug resistance mutations | # of Partially discordant mutations | # of Discordant mutations |
|-----------|----------------------|---------------|-------------------------------|----------------|--------------------------------|------------------------------------|-------------------------|
| R1        | 1,00,000             | B             | 100.00 ±0.00%                 |                | 0                              | 0                                  | 0                       |
| R2        | 12,715               | C             | 99.72 ±0.20%                  |                | 2                              | 2                                  | 0                       |
| R3        | 2,808                | C             | 99.88 ±0.05%                  |                | 3                              | 3                                  | 3                       |
| R4        | 6,65,000             | A             | 99.87 ±0.10%                  |                | 12                             | 12                                 | 12                      |
| R5        | 97,189               | C             | 99.92 ±0.05%                  |                | 18                             | 18                                 | 18                      |
| P1        | 76,600               | B             | 99.97 ±0.05%                  |                | 2                              | 2                                  | 2                       |
| P2        | Not Available        | C             | 100.00 ±0.00%                 |                | 14                             | 14                                 | 14                      |
| P3        | 14,393               | C             | 99.49 ±0.40%                  |                | 3                              | 3                                  | 3                       |
| P4        | 6,29,757             | C             | 99.87 ±0.11%                  |                | 8                              | 8                                  | 8                       |
| P5        | Not Available        | C             | 99.77 ±0.18%                  |                | 10                             | 10                                 | 10                      |
Discussion

In this study, the validation of an in-house method for HIV-1 drug resistance genotyping developed at the National AIDS Research Institute (ICMR), Pune, India was orchestrated. Accuracy of the in-house method was assessed by comparing the results of VQA HIV Genotypic Drug Resistance proficiency testing panel samples generated by the in-house method with the results of the ViroSeq method. The VQA HIV Genotypic Drug Resistance proficiency testing panel samples used in this study consisted of HIV-1 group M subtypes with known viral loads (viral load ranges 3000 to 60,633 copies/ml). All these samples (n = 26) were amplified and sequenced successfully by both in-house method and ViroSeq method with high nucleotide sequence identity (99.41 ± 0.46%). All the clinically relevant mutations were concordant by both methods and reproducible. Despite the minimal differences seen by the partial discordance in nucleotide base callings, the in-house method demonstrated an ability to identify clinically relevant mutations correctly (99.41 ± 0.46) when compared with the ViroSeq method.

In every case of nucleotide partial discordance, one method detected a mixture of the wild type and the mutant virus, the other method detected either the wild type or the mutant virus. We performed sequence editing for all validation samples using the minor peak default nucleotide mixture calling setting at 30% of the major peak in bidirectional sequences. The minor peaks at partial discordance were analyzed to determine the predominant virus. The sequence editing was performed using the Geneious software.

Figure 1. Phylogenetic analysis of the sequences generated during the evaluation of an in-house method. The phylogenetic tree was generated by using the PhyML software to create a maximum likelihood tree [13]. The reference sequences were obtained from the Los Alamos HIV Database (www.hiv.lanl.gov). IHDR- sequences generated by an in-house method; VSQ- sequences generated by the ViroSeq method; R1 to R5- sample used in the reproducibility study (highlighted in blue); P1 to P5- sample used in the precision study (highlighted in red); A, B, C, D, E are the replicates of the same sample.
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Figure 1.
The manpower and turnaround time for both the ViroSeq and the in-house methods were similar. The cost per test incurred for the in-house method (US 1128) was approximately 50% of the cost incurred using the ViroSeq method (US 3008).

In conclusion, we have evaluated a cost effective in-house method for HIV-1 drug resistance testing using plasma samples. The validated in-house method was broadly sensitive in genotyping multiple HIV-1 group M subtypes. The validation analyses indicate 100% amplification sensitivity for samples >1000 HIV-1 copies/ml, and high accuracy (99.41 ± 0.46) when compared with the ViroSeq method. In the present study, high degree of reproducibility (99.88 ± 0.10) and precision (99.82 ± 0.20) were also observed with the in-house method. The validated in-house method may be used to effectively monitor patients failing ARV therapy, as well as to collect surveillance data on the emergence and transmission of HIV-1 drug resistance isolates in resource limited countries. Moreover, the wide applications of a cost effective and validated in-house method for HIV-1 drug resistance testing will facilitate the decision making for the appropriate management of HIV infected patients and thereby reduce the risk of onset of further drug resistance-related mutations.

Supporting Information

Table S1 Supporting Information for the amplification sensitivity of an in-house method using known viral copy number samples. (DOCX)

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Author Contributions

Conceived and designed the experiments: DNC RSP SPT. Performed the experiments: DNC. Analyzed the data: DNC APN RSP SPT. Wrote the paper: DNC APN RSP SPT.
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