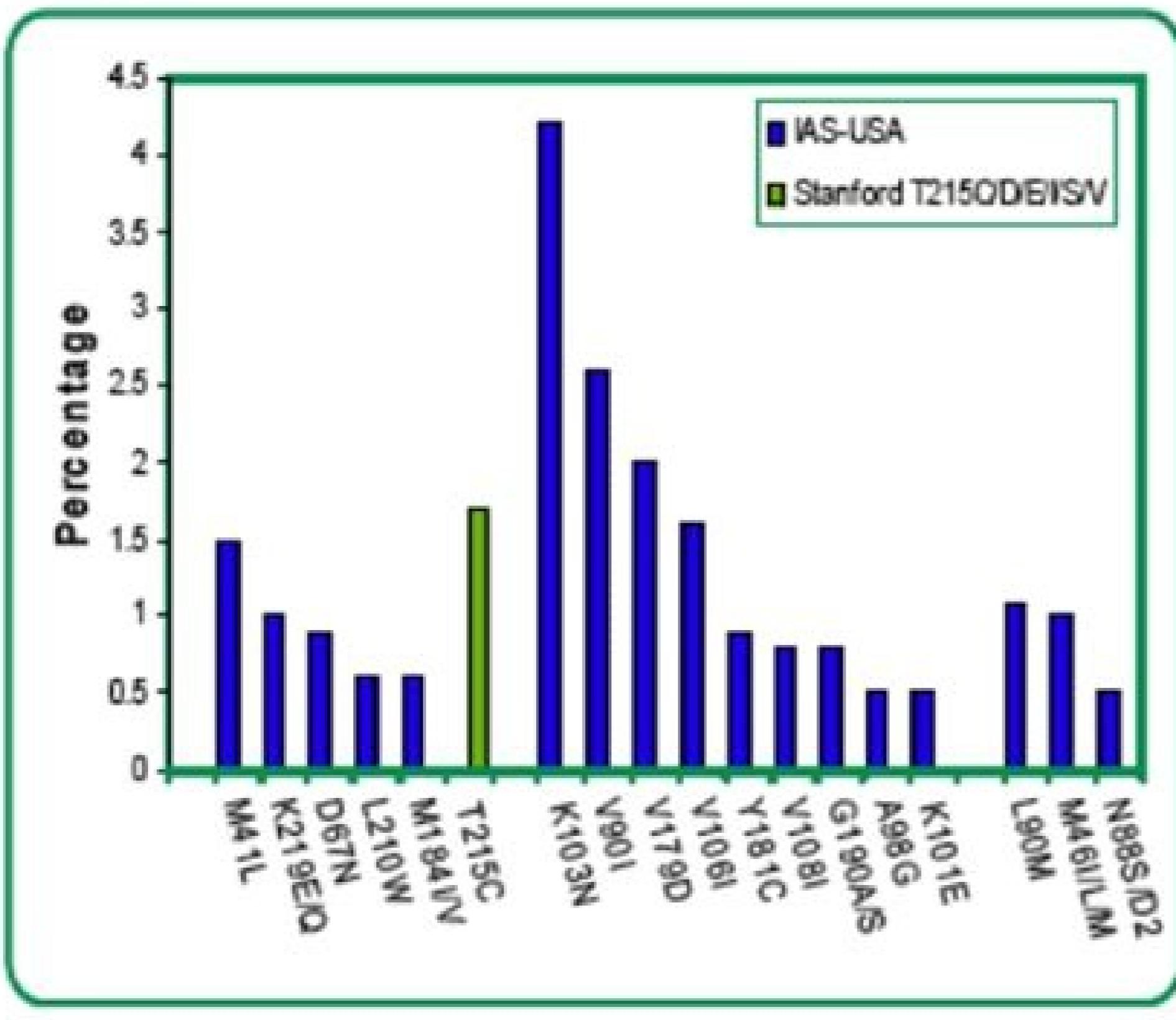


Open

Ias- usa resistance guidelines

Table 4. Advantages and Disadvantages of Currently Available Integrase Strand Transfer Inhibitors		
	2011	2012
Advantages		
Superior to nucleoside and non-nucleoside reverse transcriptase inhibitors in part due to lack of mutations associated with resistance.	Superior to nucleoside and non-nucleoside reverse transcriptase inhibitors in part due to lack of mutations associated with resistance.	Superior to nucleoside and non-nucleoside reverse transcriptase inhibitors in part due to lack of mutations associated with resistance.
Superior to nucleoside and non-nucleoside reverse transcriptase inhibitors in part due to lack of mutations associated with resistance.	Superior to nucleoside and non-nucleoside reverse transcriptase inhibitors in part due to lack of mutations associated with resistance.	Superior to nucleoside and non-nucleoside reverse transcriptase inhibitors in part due to lack of mutations associated with resistance.
Superior to nucleoside and non-nucleoside reverse transcriptase inhibitors in part due to lack of mutations associated with resistance.	Superior to nucleoside and non-nucleoside reverse transcriptase inhibitors in part due to lack of mutations associated with resistance.	Superior to nucleoside and non-nucleoside reverse transcriptase inhibitors in part due to lack of mutations associated with resistance.
Superior to nucleoside and non-nucleoside reverse transcriptase inhibitors in part due to lack of mutations associated with resistance.	Superior to nucleoside and non-nucleoside reverse transcriptase inhibitors in part due to lack of mutations associated with resistance.	Superior to nucleoside and non-nucleoside reverse transcriptase inhibitors in part due to lack of mutations associated with resistance.
Can be used with other nucleoside and non-nucleoside reverse transcriptase inhibitors.	Can be used with other nucleoside and non-nucleoside reverse transcriptase inhibitors.	Can be used with other nucleoside and non-nucleoside reverse transcriptase inhibitors.
Requires administration less frequently than nucleoside and non-nucleoside reverse transcriptase inhibitors.	Requires administration less frequently than nucleoside and non-nucleoside reverse transcriptase inhibitors.	Requires administration less frequently than nucleoside and non-nucleoside reverse transcriptase inhibitors.
Relative cost comparable to nucleoside and non-nucleoside reverse transcriptase inhibitors.	Relative cost comparable to nucleoside and non-nucleoside reverse transcriptase inhibitors.	Relative cost comparable to nucleoside and non-nucleoside reverse transcriptase inhibitors.
Can be used in patients who are unable to tolerate nucleoside and non-nucleoside reverse transcriptase inhibitors.	Can be used in patients who are unable to tolerate nucleoside and non-nucleoside reverse transcriptase inhibitors.	Can be used in patients who are unable to tolerate nucleoside and non-nucleoside reverse transcriptase inhibitors.
Highly potent.	Highly potent.	Highly potent.
Disadvantages		
Requires administration less frequently than nucleoside and non-nucleoside reverse transcriptase inhibitors.	Requires administration less frequently than nucleoside and non-nucleoside reverse transcriptase inhibitors.	Requires administration less frequently than nucleoside and non-nucleoside reverse transcriptase inhibitors.
Relative cost comparable to nucleoside and non-nucleoside reverse transcriptase inhibitors.	Relative cost comparable to nucleoside and non-nucleoside reverse transcriptase inhibitors.	Relative cost comparable to nucleoside and non-nucleoside reverse transcriptase inhibitors.
Can be used in patients who are unable to tolerate nucleoside and non-nucleoside reverse transcriptase inhibitors.	Can be used in patients who are unable to tolerate nucleoside and non-nucleoside reverse transcriptase inhibitors.	Can be used in patients who are unable to tolerate nucleoside and non-nucleoside reverse transcriptase inhibitors.
Highly potent.	Highly potent.	Highly potent.

Figure 4. Overall (2000-2007) Prevalence of Specific Resistance Mutations Observed in HIV from 15 or More Subjects



- Note: V90I, A98G, V179D, and V108I are only on the IAS-USA list and not part of the Stanford Surveillance List (version 1);
- N88S is a Stanford mutation only, but N88S alone had < 15 subjects

Evolution of DHHS Treatment Initiation Guidelines

Factor	Recommendation					
	1998	2001/2002	2004	2007	2009/2011	2012
AIDS	Treat	Treat	Treat	Treat	Treat	Treat
CD4 cell count (cells/mm ³)	Treat <500	Treat <200 Offer <350 Indiv. >350	No change	Treat <350 Risks/benefits if >350	Treat <350 Recommend ed 350-500 >500 optional	Treatment recommended at all CD4 cell counts
Viral load (copies/ml)	>20,000	>55,000	>100,000	No specific viral load	No specific viral load	No specific viral load
Other factors where treatment is recommended for all individuals with HIV-1	Pregnancy	Pregnancy	Pregnancy	Pregnancy	Pregnancy	Pregnancy
Other factors where treatment is recommended for all individuals with HIV-1	ART is recommended for all individuals with HIV-1. Effective ART has been shown to prevent transmission of HIV from an infected mother to a child. ART should be offered to patients who are at risk for transmitting HIV-1 through sexual contact and/or drug use.	ART is recommended for all individuals with HIV-1. Effective ART has been shown to prevent transmission of HIV from an infected mother to a child. ART should be offered to patients who are at risk for transmitting HIV-1 through sexual contact and/or drug use.	ART is recommended for all individuals with HIV-1. Effective ART has been shown to prevent transmission of HIV from an infected mother to a child. ART should be offered to patients who are at risk for transmitting HIV-1 through sexual contact and/or drug use.	ART is recommended for all individuals with HIV-1. Effective ART has been shown to prevent transmission of HIV from an infected mother to a child. ART should be offered to patients who are at risk for transmitting HIV-1 through sexual contact and/or drug use.	ART is recommended for all individuals with HIV-1. Effective ART has been shown to prevent transmission of HIV from an infected mother to a child. ART should be offered to patients who are at risk for transmitting HIV-1 through sexual contact and/or drug use.	ART is recommended for all individuals with HIV-1. Effective ART has been shown to prevent transmission of HIV from an infected mother to a child. ART should be offered to patients who are at risk for transmitting HIV-1 through sexual contact and/or drug use.
Other factors where treatment is recommended for all individuals with HIV-1	-2012 DHHS Guidelines					

HIVAN, HIV-associated nephropathy.

Adapted from the Panel on Antiretroviral Guidelines for Adults and Adolescents. Guidelines for the use of antiretroviral agents in HIV-1-infected adults and adolescents. US Dept of Health and Human Services; 1998-2012.

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[12]Monitoring of STIs and HAART, in general It is essential due to the disastrous effects of drug non-compliance and suboptimal treatment. The most serious rashes are Stevens-Johnson syndrome, as well as the ³ epidermal necrolysis ⁴ xica.⁵ [7]In addition to the complications of dermatitis from the use of NNRTIs, some NNRTIs are known to cause other adverse effects. With the rise of the HIV/AIDS epidemic, many companies have created drugs to reduce the spread ³ and possibly cure this problem. Drug resistance is a serious complication that must ³ be considered and is associated with poor adherence to medication⁶ [PubMed: 11 293 802].Paik S, Sen S, Era N, Saha B, Tripathi SK. In addition, factors such as the cost of treatment, accessibility to medicines and access to adequate follow-up play an important role in the development of resistance to HIV's basic regimens. First of all, a previous history of STI hypersensitivity is a contraindication ³ its use. The drug exerts its effect through the ³ yduts yduts LEEHRAT Mehta stlser Mehta: eniduvodiz ro rivacaba OT eniduvodiz mor - phe 'dehcifts stneitap detcefni suriv cneicifiedonummi namuh ni ypareht larivorterita evitca ylhgih Tiwa detaicossa yportaoipil ni tnemevorpml meht erac gnikat naicinilc yramirp Mehta Tiwa pu-wollof esolc evah diuohs, TRAAH ylnatropmi Merom Adnan, please stneitapA A elbitpecus si epytong HIV cifices richt hchih OT gurd a OT gnichtws eruquer snoitcaer esrevda gnisuc sgurd rehto Allan a] 61 [.ycacifte nemiger lamitpo niatnam llits Adnan ybportsydopil evorpmi pleh liw rivacaba OT gnichtws taht tsegus seidts, palm enduvats yllacifceps. VIH tnemtaert Mehta ni enipariven Adhan zneriavate srotibhnni esatpicrnsart esrevre edisoeilcunnon Mehta .snoitcefini lariv Miam palm owt srotibhnni esatpicrnsart esrevre esu Mehta devorppa sah ADF Mehta, etad OT. VIH Tiwa noticefini tneitap palm laitnetop sttsixe nrecnoc nehw sixalyhpmp Erusopxe-tsop of desu neeb osla evah srotibhnnart esrevre ä.] 1 [.b sittitapeh si Suriv dnoces eht. ngaid nilc j m .erteucts

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