

Neurological Manifestations of Efavirenz Toxicity:

a three-year cross-sectional study - 41 patients from Pretoria

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Background

- Efavirenz highly effective antiretroviral drug
- Backbone of HIV treatment until recently
- Well-known neuro-psychiatric side effects
- Recently described late onset toxicity
- Limited data available

Aim

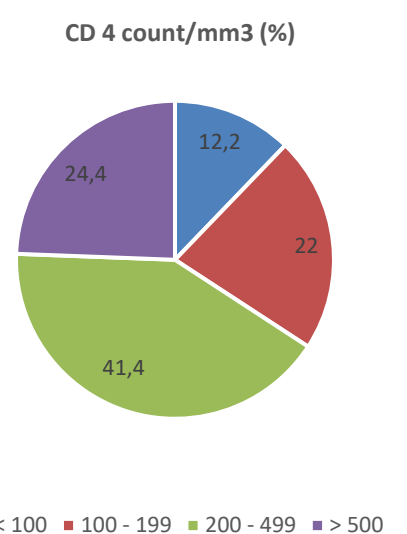
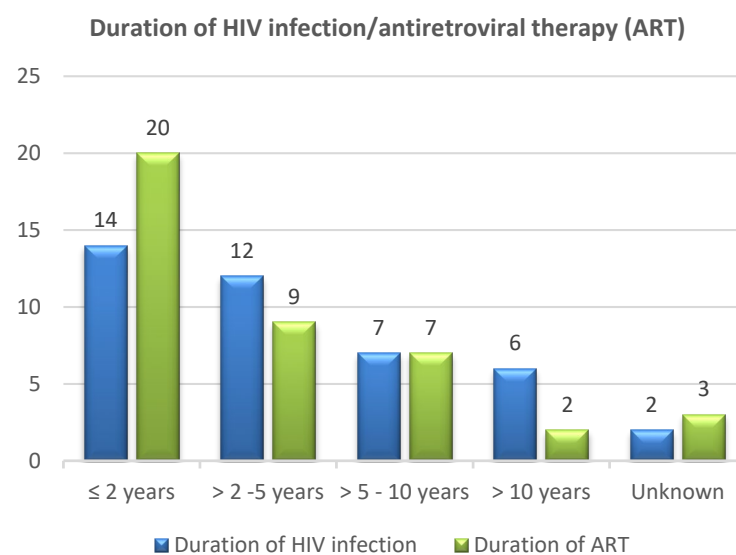
- Describe clinical picture of patients with efavirenz toxicity
- Investigate co-morbidities, risk factors
- Report outcome

Methods

- Descriptive cross-sectional study
- Three year period: 1.1.2018-31.12.2020
- Kalafong Provincial Tertiary Hospital
- All patients with efavirenz toxicity

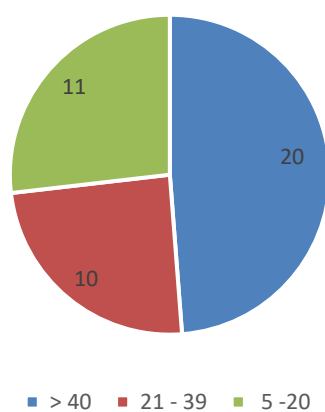
Results: HIV-related data

Variable	n=41	%
Sex		
female	36	87.8
male	5	12.2
Mean age, years (range)	43.6 (15-69)	-
female (mean)	44.75 (15-69)	
male (mean)	35.4 (17-56)	
HIV viral load (copies/ml)		
Not detectable	35	85.4
>1000	5	12.2
Unknown	1	2.4
Infection		
Tuberculosis	8	19.5
Syphilis	2	4.8
Others	2	4.8
Medication		
TB medication	9	22
INH prophylaxis	8	19.5
Sodium valproate	5	12.2

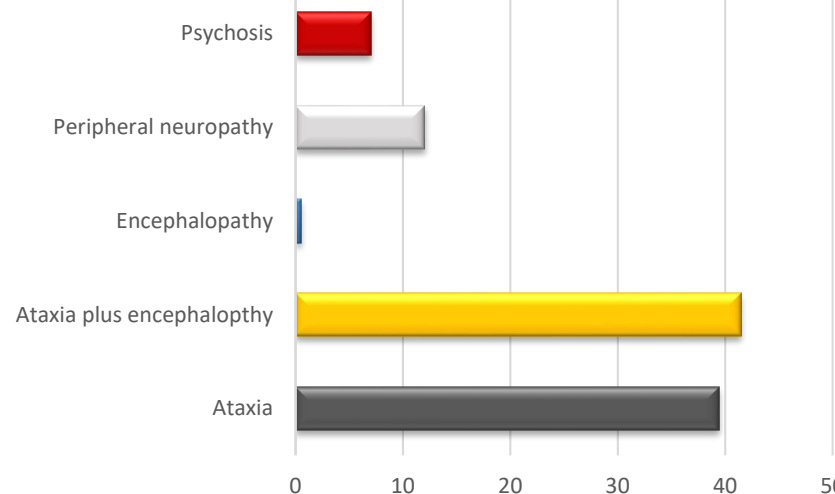


Results: Efavirenz toxicity data

Efavirenz levels µg/ml, n=41

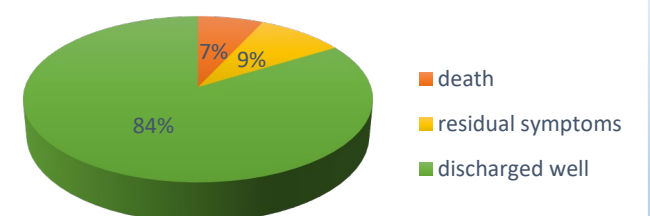


Neurological signs (%)



Efavirenz levels (n, SD)	Mean (µg/ml)	P value
Entire patient group	44.6 (41, 27.5)	-
Classical presentation	48.97 (36, 26.6)	
Psychosis/no typical symptoms	13.2 (5, 9.7)	0.0025
TB/INH treatment (Classical presentation)	52.18 (16,)	

Outcome at discharge



Conclusion

- Late onset efavirenz toxicity relatively common
- Largest single-centre cross sectional study thus far
- Ataxia and encephalopathy prominent features
- Psychosis also seen
- Prognosis good if recognized and drug withdrawn
- Mortality 7% concerning
- Risk factors: female, co-medication, especially TB treatment
- New ART guidelines replace efavirenz but many still on old regime
- Awareness of condition vital



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References

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2. Cross HM, Chetty S, Asakile MT, Hussey HS, Lee Pan EB, Tucker LM. A proposed management algorithm for late onset efavirenz neurotoxicity. S Afr Med J. 2018 Mar 28; 108(4):271-274.