



COMPARATIVE STUDY OF ANTI-HYPERTENSIVE ACTION OF AZILSARTAN WITH OLMESARTAN AND CANDESARTAN IN MILD TO MODERATE HYPERTENSION

Pharmacology

Kali Charan Rajak Tutor, Department of Pharmacology, Government Medical College Bettiah

Rahul Ranjan* Tutor, Department of Pharmacology, Government Medical College Bettiah
*Corresponding Author

Shambhu Kumar Assistance Professor, Department of Pharmacology, Government Medical College Bettiah

Awadhesh Kumar Jha Associate professor, Department of Pharmacology, Government Medical College Bettiah

ABSTRACT

Hypertension has been identified as the leading risk factor for mortality worldwide. It may lead to damage of heart, kidney, brain, vasculature and the other organs results in premature morbidity and death. The objective of the study was to compare efficacy and tolerability of once daily treatment of the new angiotensin type1 receptor blocker (ARB) Azilsartan with Olmesartan and Candesartan. **Methods:** The study was a prospective, randomized open label comparison. Total 400 patients were recruited for the study. Patients were divided into four groups. Group A comprising of 100 patients received azilsartan (40mg), Group B comprising of 100 patients received azilsartan (80mg), Group C comprising of 102 patients received olmesartan (40mg) and Group D comprising of 98 patients received candesartan (12mg). Blood pressure was monitored at base line, after 2 weeks, 4 weeks and 8 weeks of treatment. **Results:** All drugs reduced both systolic blood pressure (SBP) and Diastolic blood pressure (DSP) significantly, but the reduction in SBP and DSP with azilsartan (80mg) was significantly greater than other drugs. The difference in BP reduction between azilsartan (40mg) and olmesartan (40mg) were not significant but both azilsartan (40mg) and olmesartan (40mg) were significantly more effective than candesartan (12mg). **Conclusions:** The study indicates that azilsartan (80mg) is more effective in the control of hypertension than olmesartan and candesartan with similar safety profile.

KEYWORDS

Azilsartan, Candesartan, Essential hypertension, Olmesartan

INTRODUCTION

Hypertension is a common disorder in adults around the globe and among the most common attributable causes of mortality.[1] The goal of antihypertensive therapy is to maintain blood pressure of <140/90mmHg for most people.[2] The angiotensin receptor blockers (ARBs) have been in clinical use since 1995 and known to be effective antihypertensive agent with excellent tolerability profiles. Azilsartan medoximil, a new generation ARB for the treatment of essential hypertension. Azilsartan was discovered through the efforts of scientists from Takeda, a Japanese pharmaceutical company by modifying the tetrazole ring present in candesartan. The chemical structure of azilsartan is very similar to the structure of candesartan and differs only by replacement of candesartan's [3] member tetrazole ring with the oxa-oxadiazole ring of azilsartan. This modification makes azilsartan less acidic and more lipophilic than candesartan. Azilsartan was recently approved and has been shown to provide a more potent and sustained antihypertensive effects than other ARBs. Azilsartan medoximil, olmesartan medoximil and candesartan cilexetil are prodrugs and require activation in liver in their active forms azilsartan, olmesartan and candesartan respectively.[4,5] Molecular interaction of azilsartan with the AT(1) receptor and its strong inverse agonist activity towards the production of inositol phosphate(IP) could explain its strong BP lowering activity.

METHODS

Study design

We undertook randomized, open label comparative study of hypertensive patients in Government Medical college and Hospital, Bettiah between May 2019 to Feb 2020. Total four hundred patients were recruited for this study. Patients were randomly divided into four groups. Group A comprising of 100 patients received azilsartan (40mg), Group B comprising of 100 patients received azilsartan (80mg), Group C comprising of 102 patients received olmesartan (40mg), Group D comprising of 98 patients received candesartan (12mg) respectively.

INCLUSION CRITERIA

- Male and female of age between 25yrs to 55yrs.
- Systolic B.P between 130-169mm Hg and diastolic BP between 90-109mmHg

EXCLUSION CRITERIA

- Pregnant and lactating women
- Patients already on other antihypertensive drugs
- Patients with other condition like severe hypertension, diabetic,

hepatic failure, renal failure, heart failure, acute severe asthma

- Secondary hypertension
- Chronic use of corticosteroids, NSAIDs and sex hormones like oral contraceptive pills.

A physical examination, 12 lead electrocardiography and laboratory test were performed. Sitting cuff blood pressure was measured with mercury sphygmomanometer. Patients were seated for minimum of 5 minutes before the first measurement. Three recordings were taken, each separated by a minimum period of one minute. The pulse rate was measured once at the time of second blood pressure reading. Patients who met the entry criteria for the study during screening were assigned to receive a once daily dose of one of the following ARBs; 40mg or 80mg azilsartan, 40mg olmesartan, 12mg candesartan group wise. Patients in the treatment phase of the study were required to visit the clinic prior to taking their daily dose of medication at 2, 4 and 8 week after commencing treatment. At each visit sitting cuff blood pressure was measured in triplicate, heart rate was also measured, compliance was assessed by pill count and patients were queried for adverse events.

STATISTICAL ANALYSIS

Values are expressed as the mean $\pm$ SD. The difference of the baseline characteristics and change in BP between groups were compared using an unpaired t test. The difference between values before and after antihypertensive medication within the same group were tested using a paired t-test. P value <0.05 considered statistically significant.

RESULTS

Table 1 summarizes the baseline characteristics of the patients enrolled for this study. There were no significant differences in background factors between these groups.

The difference in blood pressure reduction after treatment with azilsartan, olmesartan and candesartan were apparent within 2 weeks. The difference in both DBP and SBP response between azilsartan (80mg) and the comparison drugs were significant for all comparisons at both 2 and 4 weeks. The difference in BP response with azilsartan (40mg) were comparable with olmesartan (40mg). Compare to candesartan (12mg), the change in BP were significant with both azilsartan (40mg) and olmesartan (40mg) (Table 2).

Table 1: Base line demographic characteristics of hypertensive patients enrolled for study.

Azilsartan (40mg)	Azilsartan (80mg)	Olmesartan (40mg)	Candesartan (12mg)
-------------------	-------------------	-------------------	--------------------

Group-A	Group-B	Group-C	Group-D		
No of patients	100	100	102	98	
Age	52±9.5	51±8.40	52±8.2	51±8.44	
Gender	Male- 67.4	Male- 66.3	Male- 67.7	Male- 68.2	
Female- 33.6	Female- 37.7	Female- 34.3	Female- 38.8		
BMI(Kg/m ²)	25±2.4	24±2.9	25±2	24±3	
Baseline blood pressure	DBP	102±3.6	102±2.7	102±2.5	102±3.4
SBP	157±12.5	158±11.6	157±1	156±12.8	

Table 2: Change in Diastolic blood pressure (ADBP) and Systolic blood pressure (ASBP) after 2 and 4 week of treatment.

Azilsartan (40mg)	Azilsartan (80mg)	Olmesartan (40mg)	Candesartan (12mg)
2 weeks	Δ DBP	-10.8	-12.7 -10.6 -9
Δ SBP	-13.7	-15.8	-13.4 -9.4
4 weeks	Δ DBP	-11.3	-14.3 -11.4 -9.7
Δ SBP	-13.8	-16.2	-13.6 -10.4

Table 3: Adverse events during the treatment period.

	Atzilsartan (40mg)	Azilsartan (80mg)	Olmesartan (40mg)	Candesartan (12mg)
Serious AEs, N (%)	0	0	0	0
Common AEs, N (%)	Headache	10(9.5)	8(7.5)	10(9.8) 9(9.1)
Dyslipidemia	9(5.7)	5(3.7)	2(1.9)	2(2.04)
Dizziness	8(5.7)	7(7.5)	5(5.8)	3(4.08)
Diarrhoea	7(1.9)	0	1(0.98)	0
Coughing	1	2(1.8)	2(1.9)	1(1.02)
Arthralgia	4(1.9)	2(1.8)	3(3.9)	6(6.12)
N=100	N=100	N=102	N=98	

DISCUSSION

Although several previous head to head comparisons of ARBs in which clinical blood pressure was used as the primary efficacy variable have been published.[6-7] Azilsartan, an angiotensin type 1 (AT1) receptor blocker (ARB) was recently approved by regulatory clinical market. The development of AT1 receptor blockers (ARBs) can be traced back to the pioneer work of scientist at Takeda pharmaceutical who described a series of benzylimidazole compounds that inhibited the ability of angiotensin to stimulate the vascular contraction and increase blood pressure (BP).[8-9] More than 15 years after the clinical introduction of Losartan, the FDA approved Takeda's azilsartan medoxomil as the 8th ARB for the treatment of hypertension.20 Azilsartan was discovered by modifying the tetrazole ring present in candesartan.[10,11] Chemical structure of azilsartan is very similar to the structure of candesartan and differ only by replacement of candesartan's 5 member tetrazole ring with the 5 member oxadiazole ring of azilsartan. Unlike candesartan which must be orally administered as a prodrug candesartan cilexetil to ensure adequate bioavailability, azilsartan has been shown to be effective in reducing BP when orally administered as either the ester prodrug, azilsartan medoxomil or as the primary compound.[3-5] During gastrointestinal absorption, azilsartan medoximidil is rapidly hydrolyzed to azilsartan, the bioactive molecule that selectively and competitively blocks angiotensin induced activation of AT1 receptor in an insurmountable fashion.[6,7] Azilsartan in clinically approved doses as azilsartan medoxomil has been shown to lower 24-hour BP in hypertensive patients significantly more than the maximum approved dose of olmesartan medoxomil, the later being considered by some to be one of the most potent ARBs for lowering BP.[8-10] Given the close structural relationship between azilsartan and candesartan, head to head studies comparing the BP effects of these two drugs are of particular interest. Azilsartan 40-80mg per day lowered systolic and diastolic BP significantly more than candesartan cilexetil (12mg).[11] The result regarding the binding affinity of azilsartan and candesartan demonstrated that these ARBs interact with the same sites in the AT1 receptor [(Tyr (113), Lys (199), and Gln (257)] The hydrogen bonding between the oxadiazole of azilsartan- Gln (257) is stronger than that between the tetrazole of candesartan-Gln (257).[9,10] An examination of the inhibition of inositol phosphate (IP) production by ARBs using constitutively active mutant receptors indicated that inverse agonist

activity required azilsartan-Gln (257) interaction and that azilsartan had a stronger activation with Gln (257) than candesartan. There was no difference among treatment groups in the incidence of clinical and laboratory adverse events. As a class, ARBs are noted for having a side effects profile similar to that of placebo.[11] A placebo group was not included in the current study, but the total adverse events rare, is similar to that reported for the placebo group in several placebo controlled trials carried out in hypertensive patients.

CONCLUSION

This study has shown that azilsartan (80mg) lowered BP to a significantly greater extent than olmesartan (40mg) and candesartan (12mg). azilsartan (40mg) was non-inferior to olmesartan (40mg). Both azilsartan (40mg) and olmesartan (40mg) are significantly more effective than candesartan (12mg). Azilsartan had a similar safety and tolerability profile to olmesartan and candesartan.

REFERENCES

1. Data from the National Health and nutrition examination survey (NHANES). National centre for health statistics. Health, United States; 2013:34.
2. Coubian AV, Bakris GL, Black HR. The seventh report of joint National Committee on prevention, Detection, Evaluation and Treatment of high blood pressure. JAMA. 2003;289:2560-723.
3. James PA, Oparil S, Carter BL, Cushman WC, Dennison-Himmelfarb C, Handler J, et al. Evidence-based guideline for the management of high blood pressure in adults: Report from the panel members appointed to the Eighth Joint National Committee (JNC 8). JAMA. 2014 Feb;311(5):507-20.
4. Weber MA, Schiffrin EL, White WB, Mann S, Lindholm LH, Kenerson JG, et al. Clinical practice guidelines for the management of hypertension in the community. The J of Clin Hyp. 2014 Jan;16(1):14-26.
5. Mancia G, Fagard R, Narkiewicz K, Redon J, Zanchetti A, Böhm M, et al. ESH/ESC guidelines for the management of arterial hypertension: The Task Force for the Management of Arterial Hypertension of the European Society of Hypertension (ESH) and of the European Society of Cardiology (ESC). Blood pressure. 2013 Aug 1;22(4):193-278.
6. Dasgupta K, Quinn RR, Zamke KB, Rabi DM, Ravani P, Daskalopoulou SS, et al. The 2014 Canadian Hypertension Education Program recommendations for blood pressure measurement, diagnosis, assessment of risk, prevention, and treatment of hypertension. Canadian Journal of Cardiology. 2014 May;30(5):485-501.
7. American Diabetic Association. Standard of medicine in diabetes and hypertension. Diabetes care. 2013;36:19-25.
8. Hübner R, Högemann AM, Sunzel M, Riddell JG. Pharmacokinetics of candesartan after single and repeated doses of candesartan cilexetil in young and elderly healthy volunteers. J of Human Hyp. 2017 Sep 2;11.
9. Scott LJ, Mc Corn PL. Olmesartan medoximil. A review of its use in the management of hypertension. Drugs. 2008;68:1239-72.
10. Kassler-Taub K, Littlejohn T, Elliott W, Ruddy T, Adler E. Comparative efficacy of two angiotensin II receptor antagonists, irbesartan and losartan, in mild-to-moderate hypertension. Am J of hyp. 2018 Apr;11(4):445-53.
11. Hedner T, Oparil S. A comparison of angiotensin antagonists Valsartan and Losartan in the treatment of essential hypertension. Am J Hypertens. 2020;12:414-7.