

Open-Label, Phase I, Pharmacokinetic Studies in Healthy Chinese Subjects to Evaluate the Bioequivalence and Food Effect of a Novel Formulation of Abiraterone Acetate Tablets

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Purpose: Abiraterone acetate tablets (I)(N-AbA) is a novel tablet co-formulated with the absorption enhancer sodium N-(8-[2-hydroxybenzoyl] amino) caprylate (SNAC). This study aimed to compare the pharmacokinetics, bioequivalence, safety, and food effects of N-AbA with the reference ZYTIGA® (R-AbA) in healthy Chinese male subjects.

Patients and Methods: This study was conducted in three parts. Part I was an open, dose-escalation trial conducted in 16 Chinese healthy males; Part II was a randomized, open-label, 2 × 4 crossover, single-dose bioequivalence trial conducted in 36 subjects; Part III was a randomized, 3 × 3 crossover trial conducted on 24 volunteers to investigate the effect of food on the pharmacokinetics of N-AbA.

Results: The exposure ($AUC_{0-\infty}$) and maximum concentration (C_{max}) of abiraterone and excipient SNAC were linear in the range of 75–450 mg dose. The bioavailability of N-AbA 300 mg was equivalent to that of R-AbA 1000 mg. The drug exposure of prednisone and prednisolone was not affected by SNAC co-administration. The C_{max} of orally administered abiraterone as R-AbA in a modified fed state was 5.9 times and $AUC_{0-\infty}$ was 4.3 times, respectively, higher than those in of orally administered abiraterone as N-AbA in a high-fat diet. The C_{max} and $AUC_{0-\infty}$ of orally administered abiraterone as N-AbA on a high-fat diet were 2.2 times and 2 times, respectively, higher than those on a fasting state. All adverse events reported in the three parts of the study were grade 1 or 2, and no serious adverse events were reported.

Conclusion: These three Phase I trials showed that N-AbA and excipient SNAC had excellent linear pharmacokinetic characteristics. A single dose of N-AbA 300 mg was bioequivalent to R-AbA 1000 mg in healthy subjects under fasting conditions. Meanwhile, SNAC had no effect on the pharmacokinetics of prednisone and prednisolone. The effect of food on N-AbA was significantly lower than that on R-AbA.

Keywords: food effect, pharmacokinetics, abiraterone acetate, bioequivalence

Introduction

Prostate cancer is one of the most common cancers among men and is the second leading cause of cancer-related death in men in the United States.¹ In Asians too, the age-adjusted incidence of prostate cancer has increased gradually over time. Androgen deprivation therapy is considered the primary treatment option for metastatic prostate cancer.² Abiraterone is a potent and selective irreversible inhibitor of the enzyme

CYP17A1, which is required for androgen biosynthesis. It slows the progression of prostate cancer by reducing the production of androgens in the testes, adrenal glands, and prostate tumor tissue.^{3,4} Abiraterone acetate, a prodrug of abiraterone and marketed under the original brand name R-AbA, is approved as a medication co-administered with prednisone for the treatment of metastatic castration-resistant prostate cancer (mCRPC) and metastatic high-risk castration-sensitive prostate cancer.⁵

Abiraterone acetate is a Biopharmaceutics Classification System class IV compound, with an extremely low oral bioavailability when administered in the fasted state.⁶ However, the administration of abiraterone acetate with high-fat food results in up to a 10-fold increased area under the concentration–time curve (AUC) and up to a 17-fold increase in the maximum plasma concentration (C_{max}) of abiraterone.⁷ Because of the low oral bioavailability and significant food effect of abiraterone acetate, its administration carries a risk of toxicity and inaccurate dosing.⁸ The specification for ZYTIGA[®] is that abiraterone acetate should be taken on an empty stomach, at least 1 h before or at least 2 h after a meal. Its large typical daily dose (1000 mg) under the fasting condition is a significant pill burden for patients.

Abiraterone acetate tablets (I)(N-AbA), developed by Jiangsu Hengrui Medicine Co., Ltd., are prepared using nanocrystalline technology in which abiraterone acetate is co-formulated with the absorption enhancer sodium N-(8-[2-hydroxybenzoyl] amino) caprylate (SNAC). Despite not being included in the FDA list of excipients, SNAC has been widely used to enhance the oral absorption of different types of drugs. The glucagon-like peptide-1 receptor agonist semaglutide co-formulated with SNAC exhibited improved drug absorption.⁹ This new co-formulation has been approved by the FDA.¹⁰ The ability of SNAC to improve drug absorption has been demonstrated in many studies.^{11,12}

N-AbA was developed to promote the gastrointestinal absorption, low oral bioavailability, high pharmacokinetic variability, and food effects of abiraterone. Here, we conducted this study aimed at comparing the pharmacokinetic characteristics, safety, and food effects of N-AbA and R-AbA.

Materials and Methods

Ethics Statement

This series of clinical trials is registered with the Chinese Clinical Trial Registry (<http://www.chictr.org.cn>;

ChiCTR2000032273, ChiCTR1900025101, ChiCTR1900025027; date of registration: April 13, 2020; August 11, 2019; April 8, 2019) and has been approved by the Institutional Review Board of The Third Xiangya Hospital of Central South University (2019070, 2019091, 2019195). The study execution dates for these trials were April 8, 2020 to May 2, 2020; July 15, 2020 to August 1, 2020; and August 29, 2020 to September 22, 2020. The study was conducted in accordance with the Declaration of Helsinki (1989), the Guidelines of Good Clinical Practice, and local applicable laws and regulations. All subjects have provided written informed consent forms prior to their participation in the study.

Study Design

This study was conducted in three parts. Part I was a single-center, open, and sequential trial conducted in 16 healthy male Chinese subjects under fasting condition. All subjects were randomized to receive a single dose of 75 mg, 150 mg, 300 mg, and 450 mg N-AbA. Each of the four doses was administered with a 7-day washout period in between each dose.

Part II was a randomized, open-label, two-sequence, four-period, single-dose bioequivalence trial conducted in 36 healthy male Chinese subjects under fasting condition. All subjects were assigned with randomized numbers into two treatment sequence groups: TRTR and RTTR (where T refers to 300 mg N-AbA with 5 mg prednisone, and R refers to R-AbA with 5 mg prednisone). The subjects received T or R products according to the randomization schedule. The washout period was 7 days.

Part III was a randomized, open-label, three-sequence, three-period, self-crossover study conducted in 24 healthy male Chinese subjects to compare the food effects on pharmacokinetic and safety profiles between the two formulations. All subjects were randomized into one of the three cohorts: 1) subjects were administered 300 mg N-AbA under fasting state; 2) subjects were administered 300 mg N-AbA on a high-fat diet; 3) subjects were administered 1000 mg R-AbA on modified fed conditions. The washout period was 7 days.

For the fasting condition, the subjects were asked to fast overnight (10 h) prior to dosing. The high-fat diet contained about 800–1000 kcal in total with about 50% of the calories coming from fat. The high-fat meal was provided 30 min before administration and finished within 30 min. Under the modified fed condition, subjects were asked to eat a medium-fat meal (~450 kcal with 30% of the calories coming from fat) 2 h before and 1 h after

administration. No water was consumed between 1 h before and 1 h after drug administration, except the 240 mL water to swallow the study drugs.

Study Drugs

The test drug (N-AbA) was manufactured by Chengdu Suncadia Medicine Co., Ltd. (subsidiary of Jiangsu Hengrui Medicine Co., Ltd.); specifications: 150 mg per tablet; batch number: P20060411. The reference drug (R-AbA) was manufactured by Patheon Inc.; specifications: 250 mg per tablet; batch number: CCTSV. The combination drug prednisone acetate tablets were manufactured by Shanghai Pharmaceuticals Sine Co., Ltd.; specifications: 5 mg per tablet; batch number: 018190903.

Pharmacokinetic Evaluation, Blood Sample Collection, and Bioanalytical Assay

Plasma concentrations of abiraterone and SNAC were measured in all three trials in this study. In Part II, prednisone and prednisolone concentrations were also tested. Blood samples were collected prior to dosing (0 h) and at 0.25, 0.5, 0.75, 1, 1.25, 1.5, 1.75, 2, 2.5, 3, 4, 5, 6, 8, 12, 24, 36 and 48 h post-dose. For Part II, blood samples were also collected at 3.5 and 4.5 h post-dose. Blood samples were centrifuged at $1300 \times g$ for 10 min at 4°C. The obtained plasma samples were stored in a refrigerator at $-20 \pm 5^\circ\text{C}$ for pre-freezing, and the samples were transferred to the refrigerator at $\leq -60^\circ\text{C}$ on the last day in each period for subsequent analyses. The time from the end of centrifugation to freezer storage was less than 1 h.

The serum concentration of abiraterone, SNAC, prednisone, and prednisolone was detected by HPLC-MS/MS. The limit of quantification of abiraterone, SNAC, prednisone, and prednisolone was 0.100–400, 0.250–1000, 0.500–50.0, and 2.00–200 ng/mL. Of all analytical methods of four compounds, the percent co-efficient of variation of inter-assay precision was $\leq 7.4\%$, and the assay accuracy was in the range of -8.0% to 1.7% .

Safety Assessments

Safety was evaluated by observing vital signs, physical examination, electrocardiography, laboratory examination (blood routine, blood biochemistry, blood coagulation function, urine routine, fecal routine, fecal occult blood, and blood transfusion), adverse event (AE) reporting. AEs were recorded and graded according to the National

Cancer Institute Common Terminology Criteria for Adverse Events (version 5.0). Subjects with AEs were followed up until the symptoms or the corresponding physical and clinical examinations returned to normal.

Statistical Analysis

Pharmacokinetic parameters were calculated using the noncompartmental model (NCA module) on WinNonlin 8.2 (Pharsight Corporation, Mountain View, CA, USA). Statistical analysis was performed with SAS 9.4 Statistical Package (SAS Institute Inc., Cary, NC, USA). In Part I, the confidence interval method was used to analyze the linear relationship of AUC and C_{\max} with the dose. The natural logarithm of AUC and C_{\max} was used to perform linear regression analysis with the dose. The slope of the linear regression equation and 90% confidence interval were observed. In Part II, natural log-transformed AUC_{0-t} , $\text{AUC}_{0-\infty}$, and C_{\max} were analyzed by analysis of variance with a mixed-effects model. Treatment sequence, formulation, and period were the fixed effect variables, and participant was the random effect variable. The relative bioavailability of the test drug and reference drug was evaluated based on average bioequivalence (ABE) or reference-scaled average bioequivalence (RSABE). In Part III, the least-squares mean ratios and 90% confidence intervals of C_{\max} , AUC_{0-t} , and $\text{AUC}_{0-\infty}$ of abiraterone in the plasma were calculated after oral administration under different feeding conditions. The adjusted mean differences and 90% CIs for the differences were exponentiated to provide estimates of the ratio of adjusted geometric means (test/reference) and 90% CIs for the ratios.

Results

Participant Demographics

A total of 76 healthy male subjects were enrolled in this study (Part I: $n = 16$; Part II: $n = 36$; Part III: $n = 24$). One subject dropped out in each of the three parts of the study due to AEs. The flowchart of the subject distribution is shown in Figure 1. The median age of the subjects was 28.25, 27.81, and 25.08 years in the three trials. The median body mass index was 21.61, 22.24, and 21.44 kg/m². The participant characteristics are shown in Table 1.

Pharmacokinetics

Part I

The plasma concentration profiles and pharmacokinetic parameters of abiraterone and SNAC are shown in

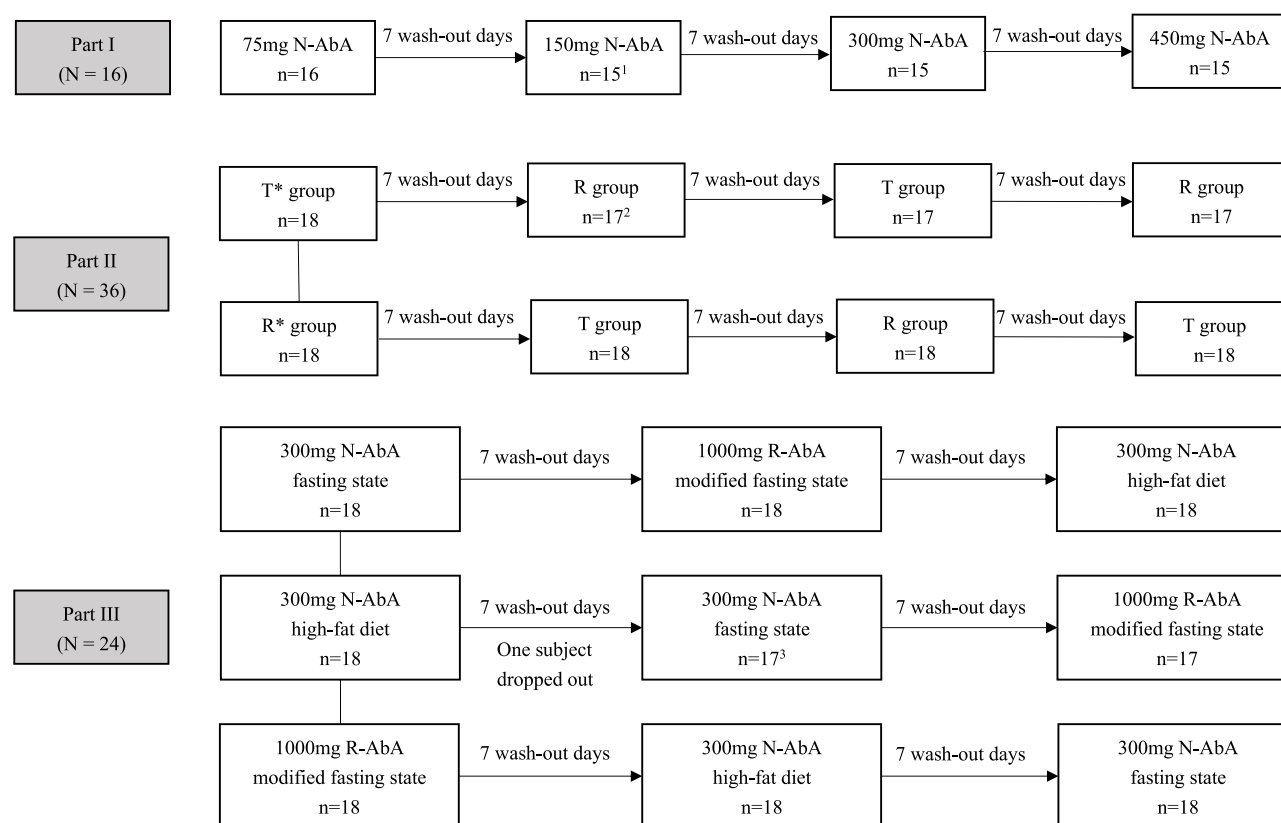


Figure 1 The flowchart of the subject disposition. ¹one subject dropped out because of elevating the level of uric acid. ²one subject dropped out because of elevating the level of triglycerides. ³one subject dropped out because of elevating the level of blood bilirubin. *T group: 300mg N-AbA + 5mg prednisone; 1000mg R-AbA + 5mg prednisone.

Table 2. The median T_{max} of abiraterone and SNAC was approximately 1 h and 0.5 h, respectively. Following the oral administration of 75–450 mg AbA, the C_{max} and

$AUC_{0-\infty}$ of abiraterone and SNAC were proportional to the dose (Figure 2).

Part II

After the administration of 300 mg N-AbA with 5 mg prednisone tablet or 1000 mg R-AbA with 5 mg prednisone tablet, the mean plasma concentration profiles under fasting condition are shown in Figure 3. For the test group, compared to the reference group, the geometric mean ratios of C_{max} , $AUC_{0-\infty}$, and AUC_{0-t} were 106.62%, 92.38%, and 94.01%, respectively. All 90% confidence intervals (CIs) were within the range of 80–125%. To evaluate the effect of SNAC on the pharmacokinetics of prednisone, we compared the C_{max} , $AUC_{0-\infty}$, and AUC_{0-t} of prednisone and prednisolone between the two groups and found that these pharmacokinetic parameters of prednisone and prednisolone were similar between the test and reference groups (Table 3).

Part III

The pharmacokinetic profiles of abiraterone under fasting conditions (300 mg N-AbA), high-fat diet conditions

Table 1 Demographics and Characteristics of Subjects

Characteristics	Part I (n=16)	Part II (n=36)	Part III (n=24)
Age (year)			
Median±SD	28.25±6.34	27.81±6.15	25.08±5.93
Range	19.00~37.00	20.00~45.00	18.00~38.00
Gender (n%)			
Male	16(100.0%)	36(100.0%)	24(100.0%)
Female	0(0.0%)	0(0.0%)	0(0.0%)
Weight (kg)			
Median±SD	61.07±6.67	62.76±7.11	61.25±5.16
Range	50.60~75.90	51.50~79.20	53.70~73.10
BMI (kg/m²)			
Median±SD	21.61±1.49	22.24±1.88	21.44±1.81
Range	19.40~24.10	19.30~25.70	19.10~25.90
Height (cm)			
Median±SD	167.91±6.45	167.86±6.21	169.06±4.73
Range	156.50~178.00	149.5~178.50	160.00~176.50

Abbreviation: BMI, body mass index.

Table 2 PK Parameters of Abiraterone and SNAC After Dosing Abiraterone Acetate with Different Doses

Parameters	Abiraterone			
Mean \pm SD (CV%)	75 mg (n=16)	150 mg (n=15)	300 mg (n=15)	50 mg (n=15)
C_{max} (ng/mL)	33.9 \pm 12.9 (38.1)	81.4 \pm 45.0 (55.3)	209 \pm 142(67.8)	405 \pm 428(106)
AUC_{0-t} (ng·h/mL)	98.96 \pm 46.29 (46.77)	246.11 \pm 97.19 (39.49)	620.08 \pm 370.19 (59.70)	1041.37 \pm 903.24 (86.74)
$AUC_{0-\infty}$ (ng·h/mL)	101.27 \pm 46.61 (46.02)	250.27 \pm 97.95 (39.14)	627.60 \pm 370.44 (59.02)	1054.58 \pm 904.86 (85.80)
T_{max} *(h)	1.13 (0.5–2.5)	1.0(0.5–1.25)	1.0 (0.5–2.00)	1.0 (0.75–3.00)
$t_{1/2z}$ (h)	10.15 \pm 3.1 (30.52)	10.71 \pm 2.03 (18.96)	10.62 \pm 1.80 (16.94)	10.85 \pm 1.51 (13.95)
Parameters	SNAC			
Mean \pm SD (CV%)	75 mg (n=16)	150mg (n=15)	300 mg (n=15)	50mg (n=15)
C_{max} (ng/mL)	407 \pm 140 (34.4)	977 \pm 385 (39.4)	1870 \pm 484 (25.9)	2460 \pm 963 (39.2)
AUC_{0-t} (ng·h/mL)	291.34 \pm 61.44(21.09)	668.36 \pm 153.26 (22.93)	1396.58 \pm 284.15 (20.35)	2121.8 \pm 491.85 (23.18)
$AUC_{0-\infty}$ (ng·h/mL)	294.82 \pm 61.4 (20.83)	672.15 \pm 152.2 (22.64)	1409.79 \pm 285.05 (20.22)	2131.49 \pm 490.78 (23.03)
T_{max} *(h)	0.38 (0.25–1.25)	0.5(0.25–0.75)	0.5 (0.25–1.75)	0.5 (0.25–0.75)
$t_{1/2z}$ (h)	3.53 \pm 2.54(71.79)	2.94 \pm 1.39 (47.26)	5.35 \pm 3.92 (73.26)	4.88 \pm 2.37 (48.47)

Abbreviations: C_{max} , the highest plasma concentration of the drug; AUC_{0-t} , the area under the plasma concentration–time curve from time zero to the last quantifiable concentration time; $AUC_{0-\infty}$, the area under the plasma concentration–time curve from time zero extrapolated to infinite time.

(300 mg N-AbA), and modified fed conditions (1000 mg R-AbA) are shown in Figure 4. The C_{max} of abiraterone as orally administered R-AbA on a modified fed state was 5.89 times and $AUC_{0-\infty}$ was 4.3 times, respectively, higher

than that of abiraterone as orally administered N-AbA on a high-fat diet. Under the fasting state, the C_{max} of abiraterone as orally administered N-AbA on a high-fat diet was 2.2 times and $AUC_{0-\infty}$ was 2 times, respectively,

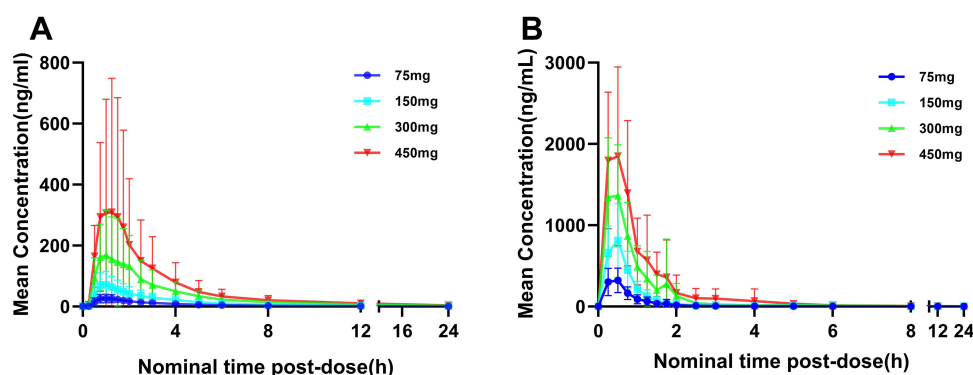
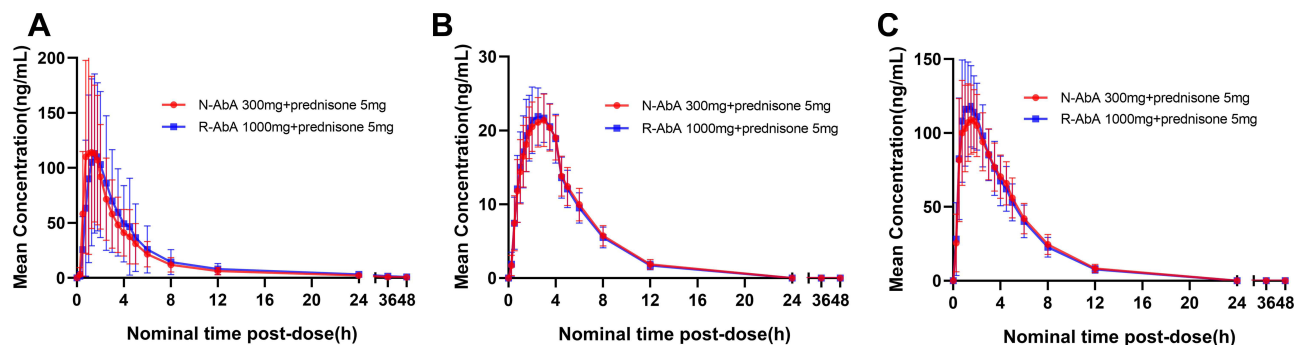
**Figure 2** Linear scale of mean plasma concentration–time plot of Abiraterone (A) and SNAC (B).**Figure 3** Mean plasma concentration–time plot of Abiraterone (A), prednisone (B) and prednisolone (C) for R and T groups.

Table 3 Summary of Bioequivalence Analysis Under Fasting Conditions

PK Parameters	Geometric Mean (Abiraterone)			90% CI
	T*(n=36)	R*(n=35)	Ratio (%)	
C _{max} (ng/mL)	151.67	142.25	106.62	95.73~118.76
AUC _{0-t} (ng·h/mL)	456.50	485.58	94.01	87.42~101.10
AUC _{0-∞} (ng·h/mL)	464.43	502.74	92.38	86.00~99.23
PK Parameters	Geometric Mean (Prednisone)			90% CI
	T*(n=36)	R*(n=35)	Ratio (%)	
C _{max} (ng/mL)	22.58	23.31	96.89	94.22%~99.63%
AUC _{0-t} (ng·h/mL)	122.16	121.88	100.23	98.26%~102.24%
AUC _{0-∞} (ng·h/mL)	128.85	127.77	100.84	98.81%~102.92%
PK Parameters	Geometric Mean (Prednisolone)			90% CI
	T*(n=36)	R*(n=35)	Ratio (%)	
C _{max} (ng/mL)	125.41	141.28	88.77	86.08%~91.54%
AUC _{0-t} (ng·h/mL)	581.96	582.63	99.89	98.29%~101.50%
AUC _{0-∞} (ng·h/mL)	611.50	609.44	100.34	98.70%~102.01%

Note: * T: 300mg N-AbA+5mg prednisone; R: 1000mg R-ABA+5mg prednisone.

higher. For SNAC, the geometric mean ratios of C_{max}, AUC_{0-t}, and AUC_{0-∞} were 71.64%, 126.17%, and 125.92%, respectively, when comparing the high-fat diet condition (300 mg N-AbA) with fasting condition (300 mg N-AbA). The details are shown in Table 4.

Safety

In Part I, 16 AEs were reported in 9 subjects (56.3%, 9/16). All AEs were of grade 1. Of these, 14 AEs were related to the study drug. There was no significant trend between dose level and incidence of AEs. One subject discontinued Part I because of increased uric acid levels. In Part II, 24 AEs occurred in 9 subjects (25%, 9/36). The frequency of AE occurrence was similar between the test group and the reference group (11 vs 13). Of those, 9 subjects reported 21 drug-related AEs (50%, 8/

16). One subject terminated the study due to elevated blood triglyceride levels. In Part III, 28 AEs occurred in 12 subjects (50%, 12/24), of which 19 AEs were related to the study drug. One subject terminated the study because of elevated levels of blood bilirubin. In all three trials, no severe AEs or AEs that caused death were reported.

The most frequent treatment-related AEs were hemobilirubin and hypertriglyceridemia. The incidence of elevated hemobilirubin and hypertriglyceridemia among patients treated with N-AbA and R-AbA did not differ. Other AEs included urinary tract infection, cough, hypertension, arrhythmia, and diarrhea. The details are shown in Table 5. All treatment-related AEs in this study are reported in the specification for ZYTIGA®.

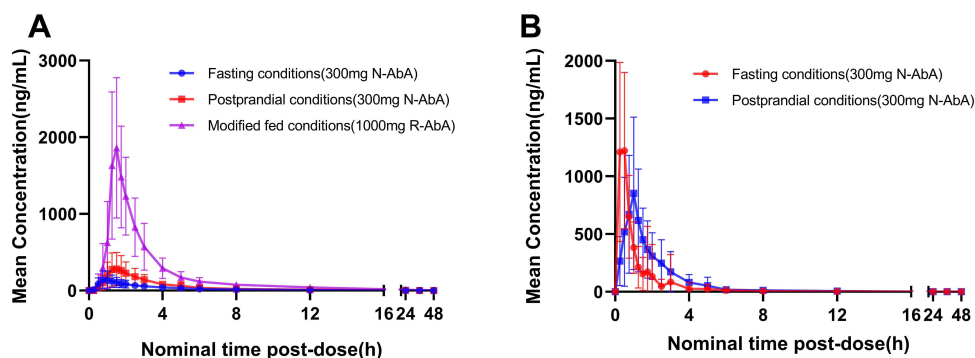


Figure 4 Mean plasma concentration–time plot of Abiraterone (A) and SNAC (B) under fasting conditions, postprandial conditions and modified fed conditions.

Table 4 Mean Pharmacokinetic Parameters of Abiraterone and SNAC

PK Parameters	Geometric Mean (Abiraterone)			90% CI
	Modified Fed Condition (1000 mg R-ABA) (n=23)	Postprandial Conditions (300 mg N-AbA) (n=24)	Ratio (%)	
C_{max} (ng/mL)	2000.25	339.77	588.71	458.06%–756.63%
AUC_{0-t} (ng·h/mL)	4037.82	938.97	430.03	368.90%–501.29%
$AUC_{0-\infty}$ (ng·h/mL)	4079.87	951.78	428.66	368.09%–499.19%
PK Parameters	Geometric Mean (SNAC)			90% CI
	Postprandial Conditions (300 mg N-ABA) (n=24)	Fasting Conditions (300 mg N-AbA) (n=23)	Ratio (%)	
C_{max} (ng/mL)	971.82	1356.56	71.64	52.71%–97.36%
AUC_{0-t} (ng·h/mL)	1523.99	1207.92	126.17	114.72%–138.75%
$AUC_{0-\infty}$ (ng·h/mL)	1531.14	1215.99	125.92	114.57%–138.39%

Discussion

The main results of Part I showed linear pharmacokinetic characteristics of abiraterone and SNAC in the range of 75–450 mg after a single oral administration of N-AbA on an empty stomach. N-AbA was safe and well tolerated during the trial. In the 300 mg cohort, the C_{max} and $AUC_{0-\infty}$ of abiraterone were 81.4 ng/mL and 250.27 ng·h/mL, respectively, which were similar to the values reported in previous studies on R-AbA.¹³ Based on these data, we further evaluated and compared the pharmacokinetics of 300 mg N-AbA and 1000 mg R-AbA.

The results of Part II showed that under fasting condition, the test/reference geometric mean ratios are all within the range of 80% to 125% indicating that the exposure of N-AbA 300 mg was bioequivalent with that of R-AbA 1000 mg, and the intra-individual variation in AUC_{0-t} and $AUC_{0-\infty}$ for N-AbA was lower than that for R-AbA. As the CYP17 inhibitor, abiraterone acetate had some side effects such as fluid retention, hypertension, and hypokalemia. Low-dose prednisone decreases steroid build-up upstream of CYP17 and prevents mineralocorticoid excess.¹⁴ In 2012, AbA plus prednisone was approved by FDA for the treatment of patients with mCRPC. European Association of Urology guidelines on the treatment of CRPC mention this combination as one of the first-line treatments for CRPC.¹⁵ Thus, in Part II, we also studied whether SNAC would affect prednisone pharmacokinetics. The pharmacokinetic parameters of prednisone and prednisolone were similar after the combined

use of N-AbA or R-AbA, indicating that SNAC had no effect on the pharmacokinetics of prednisone and prednisolone.

The specifications for ZYTIGA[®] state that the tablets must be taken on an empty stomach and should not be taken at least 2 h before and at least 1 h after administration because food effects can result in fluctuations in drug exposure. Therefore, in Part III, we set up a modified fed group based on these two time nodes. The results showed that after oral administration of the test drug after a high-fat meal, the C_{max} of abiraterone increased only by about 2.2 times, and the $AUC_{0-\infty}$ increased by about 2 times, indicating that N-AbA has a significantly lower food effect than R-AbA. Thus, even if patients do not follow regular diet management before and after taking the drug, drug exposure will not be drastically altered, which is an important safety advantage.

YONSA[®], the improved version of R-AbA, was approved for the treatment of patients with mCRPC in 2018. The proprietary YONSA[®] formulation using SoluMatrix Fine Particle TechnologyTM allows for the same systemic exposure to be achieved with 500 mg relative to the recommended dose of R-AbA 1000 mg.¹⁶ The C_{max} of abiraterone was approximately 6.5-fold higher, and the $AUC_{0-\infty}$ was 4.4-fold higher when a single dose of YONSA 500 mg was administered with a high-fat meal (56–60% fat, 900–1000 calories) than when administered after overnight fasting in healthy subjects.¹⁷ Nanocrystalline technology and the addition of SNAC further improve the bioavailability of N-AbA and reduce the impact of food.

Table 5 Summary of Treatment-Related AEs Occurred in the Study

AEs (No. of AEs, No. (%) of Subjects)	Part I (n=16)			
	75 mg (n=16)	150 mg (n=15)	300 mg (n=15)	450 mg (n=15)
Laboratory	3(3, 18.8)	6(4, 26.7)	6(6, 40.0)	1(1, 6.7)
Triglyceride increased	1(1, 6.3)	1(1, 6.7)	3(3, 20.0)	1(1, 6.7)
Bacteria	1(1, 6.3)	1(1, 6.7)	0(0, 0)	0(0, 0)
Hemobilirubin increased	0(0, 0)	2(2, 13.3)	0(0, 0)	0(0, 0)
Urokinase protein	0(0, 0)	1(1, 6.7)	0(0, 0)	0(0, 0)
Urine erythrocyte	0(0, 0)	1(1, 6.7)	0(0, 0)	0(0, 0)
Hematuresis	0(0, 0)	0(0, 0)	1(1, 6.7)	0(0, 0)
TBA increased	0(0, 0)	0(0, 0)	1(1, 6.7)	0(0, 0)
AEs (No. of AEs, No. (%) of Subjects)	Part II (n=36)			
	T*(n=36)	R*(n=35)		
Laboratory	10(7, 19.4)	13(6, 17.1)		
Triglyceride increased	5(4, 11.1)	6(3, 8.6)		
Bacteria	0(0, 0)	0(0, 0)		
Hemobilirubin increased	3(2, 5.6)	2(1, 2.9)		
DBil increased	0(0, 0)	1(1, 2.9)		
Urine leukocyte	0(0, 0)	1(1, 2.9)		
AST increased	1(1, 2.8)	0(0, 0)		
Gastrointestinal	1(1, 2.8)	0(0, 0)		
Ventosity	1(1, 2.8)	0(0, 0)		
AEs (No. of AEs, No. (%) of subjects)	Part III (n=24)			
	Fasting Conditions (T [#] , n=23)	Postprandial Conditions (T [#] , n=24)	Modified Fed Conditions (R [#] , n=23)	
Laboratory	9(7, 30.4)	9(7, 29.2)	5(3, 13.0)	
Triglyceride increased	4(4, 17.4)	4(4, 16.7)	1(1, 4.3)	
Hemobilirubin increased	0(0, 0)	1(1, 4.2)	0(0, 0)	
TBA increased	1(1, 4.3)	0(0, 0)	0(0, 0)	
WBC increased	0(0, 0)	1(1, 4.2)	0(0, 0)	
Liver enzymes increased	0(0, 0)	0(0, 0)	1(1, 4.3)	
Neutrophil count increased	0(0, 0)	1(1, 4.2)	0(0, 0)	
Gastrointestinal	0(0, 0.0%)	0(0, 0)	4(2, 8.3)	
Stomachache	0(0, 0.0%)	0(0, 0)	2(2, 8.3)	
Diarrhea	0(0, 0.0%)	0(0, 0)	2(2, 8.3)	
Respiratory	0(0, 0.0%)	0(0, 0)	2(2, 8.3)	
Cough	0(0, 0.0%)	0(0, 0)	2(2, 8.3)	

Notes: T*: 300mg N-AbA+5mg prednisone; R*: 1000mg R-AbA+5mg prednisone. T[#]: 300mg N-AbA; R[#]: 1000mg R-AbA.

Abbreviations: AE, adverse event; TBA, total bile acid; WBC, white blood cells; AST, aspartate aminotransferase.

SNAC is a new type of absorption enhancer originally developed by Emisphere Technology Inc. Despite it not being listed in the FDA excipient database at present, SNAC is widely used in improving the oral absorption of different types of drugs, including proteins and macromolecules, such as insulin, calcitonin, and heparin.^{18,19} SNAC has been reported to enhance the oral bioavailability of the

long-acting GLP-1 analog semaglutide, which is used to treat type 2 diabetes and was approved by the FDA on September 20, 2019.^{20,21} A vitamin B12 product that exhibits enhanced oral absorption due to SNAC is already marketed.²² This study confirms that the use of SNAC not only improves the bioavailability of abiraterone but also greatly reduces the effect of food on the absorption of

abiraterone. SNAC may be used to enhance the oral bioavailability of several other drugs.

Conclusion

N-AbA and the excipient SNAC showed linear pharmacokinetic characteristics within the dose range of 75 mg–450 mg. A single 300 mg dose of N-AbA was bioequivalent to 1000 mg of R-AbA in healthy subjects under fasting conditions. Meanwhile, no pharmacokinetic interactions were observed between SNAC and prednisone after combined administration of prednisone. N-AbA shows lower intra-individual variation and food effects than R-AbA.

Data Sharing Statement

The original contributions presented in the study are included in the article, and further inquiries can be directed to the corresponding author.

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Disclosure

YH was employed by Jiangsu Hengrui Medicine Co., Ltd. The authors report no other conflicts of interest in this work.

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