

## Data Sheet

# ANACCORD

### *Anastrozole*

### *1 mg tablets*

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## Presentation

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Anaccord Tablets 1mg are presented as white to off white, round, biconvex, film coated tablets containing 1 mg of anastrozole. The tablets are 6 mm in diameter and are compressed to a weight of 100 mg. A logo consisting of "AHI" debossing on one side and plain on other side of the tablets.

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## Uses

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### *Actions*

Anastrozole is a potent and highly selective non-steroidal aromatase inhibitor. In post-menopausal women, oestradiol is produced primarily from the conversion of androstenedione to oestrone through the aromatase enzyme complex in peripheral tissues. Oestrone is subsequently converted to oestradiol. Reducing circulating oestradiol levels has been shown to produce a beneficial effect in women with breast cancer. In post-menopausal women, anastrozole at a daily dose of 1 mg produced oestradiol suppression of greater than 80% using a highly sensitive assay.

In clinical trials treatment with anastrozole at a dose of 1 mg has demonstrated significant prolongation of survival time.

Anastrozole does not possess any progestogenic, androgenic or oestrogenic activity.

Daily doses of anastrozole up to 10 mg do not have any effect on cortisol or aldosterone secretion, measured before or after standard ACTH challenge testing. Corticoid supplements are therefore not needed.

Extensive phase III clinical study programmes showed that anastrozole is an effective treatment of early breast cancer and advanced breast cancer in postmenopausal women suitable for endocrine therapy.

### Primary Adjuvant treatment of early breast cancer

In a large phase III study conducted in 9366 postmenopausal women with operable breast cancer, anastrozole was shown to be statistically superior to tamoxifen in recurrence-free survival. The incidence of contralateral breast cancer was statistically significantly reduced for anastrozole compared to tamoxifen. Time to distant recurrence was also numerically superior for anastrozole. The combination of Anastrozole and tamoxifen did not demonstrate any efficacy benefits in comparison to tamoxifen.

For the prospectively defined receptor positive population, even greater statistical superiority was observed for recurrence-free survival in favour of Anastrozole versus tamoxifen. Again, the combination of anastrozole and tamoxifen did not demonstrate any efficacy benefits in comparison with tamoxifen in this group of patients.

### **Adjuvant treatment of early breast cancer for patients being treated with adjuvant tamoxifen.**

In a phase III trial (ABCSG 8) conducted in 2579 postmenopausal women with hormone receptor positive early breast cancer being treated with adjuvant tamoxifen, patients had a superior disease-free survival when switched to anastrozole compared with those continuing on tamoxifen.

Time to any recurrence, time to local or distant recurrence and time to distant recurrence confirmed a statistical advantage for anastrozole, consistent with the results of disease free survival. The incidence of contralateral breast cancer was very low in the two treatment arms, with a numerical advantage for anastrozole. Overall survival was similar for the two treatment groups.

Two further similar trials (GABG/ARNO 95 and ITA) with anastrozole, as well as a combined analysis of ABCSG 8 and GABG/ARNO 95, supported these results.

The anastrozole safety profile in these 3 studies was consistent with the known safety profile established in post-menopausal women with hormone-receptor positive early breast cancer.

### **Study of anastrozole with the bisphosphonate risedronate (SABRE)**

#### BMD

In the phase III/IV SABRE study, 234 postmenopausal women with hormone receptor positive early breast cancer scheduled for treatment with anastrozole were stratified to low, moderate and high-risk groups according to their existing risk of fragility fracture. All patients received treatment with vitamin D and calcium. Patients in the low risk group received anastrozole alone, those in the moderate group were randomised to anastrozole plus bisphosphonate or

anastrozole plus placebo and those in the high risk group received anastrozole plus bisphosphonate.

The 12-month main analysis has shown that patients already at moderate to high risk of fragility fracture had their bone health (assessed by bone mineral density and bone formation and resorption markers) successfully managed by using anastrozole in combination with a bisphosphonate. In addition, no changes in BMD were seen in the low risk group treated with anastrozole alone and given vitamin D and calcium. These findings were mirrored in the secondary efficacy variable of change from baseline in total hip BMD at 12 months.

This study provides evidence that postmenopausal women with early breast cancer scheduled to be treated with anastrozole should have their bone status managed according to treatment guidelines already available for postmenopausal women at similar risk of fragility fracture.

#### Lipids

In the SABRE study there was a neutral effect on plasma lipids both in those patients treated with anastrozole alone and in those treated with anastrozole plus a bisphosphonate.

#### Paediatrics

Three clinical trials were conducted in paediatric patients (2 in pubertal boys with gynecomastia and 1 in paediatric girls with McCune Albright Syndrome).

#### **Gynecomastia Study**

Trial 0006 was a randomised, double-blind, multi-centre study, of 80 pubertal boys with gynaecomastia of greater than 12 months duration (aged 11-18 years inclusive) treated with anastrozole 1 mg/day or placebo for up to 6 months. A decrease of  $\geq 50\%$  in total breast volume measured by ultrasound was seen in 38.5% (15/39) of the anastrozole and 31.4% (11/35) of the placebo treated group, (odds ratio = 1.513, 95% CI 0.496 to 4.844,  $p=0.4687$ ).

Trial 0001 was an open-label, multiple-dose pharmacokinetic (PK) study of anastrozole 1 mg/day in 36 pubertal boys with gynecomastia of less than 12 months duration. A decrease in total breast volume of 50% or greater at 6 months was seen in 55.6% (20/36) of the boys.

#### **McCune Albright Syndrome (MAS) Study**

Trial 0046 was an international, multi-centre, open-label, exploratory trial of anastrozole in 28 girls (aged 2 to  $\leq 10$  years) with McCune Albright Syndrome (MAS). No statistically significant change in the frequency of vaginal bleeding days on treatment was observed. Of the patients with baseline vaginal bleeding,

28% experienced a  $\geq 50\%$  reduction in the frequency of bleeding days on treatment; 40% experienced a cessation over a 6-month period, and 12% experienced a cessation over a 12-month period. There were no clinically significant changes in Tanner staging, mean ovarian volume or mean uterine volume. No statistically significant change in the rate of increase in bone age on treatment compared to the rate during baseline was observed. Growth rate (in cm/year) was significantly reduced ( $p < 0.05$ ) from pre-treatment through month 0 to month 12, and from pre-treatment to the second 6 months (month 7 to month 12).

The overall assessment of the AEs in children less than 18 years of age raised no safety and tolerability concerns.

### ***Pharmacokinetics***

Absorption of anastrozole is rapid and maximum plasma concentrations typically occur within two hours of dosing (under fasted conditions). Anastrozole is eliminated slowly with a plasma elimination half-life of 40 to 50 hours. Food slightly decreases the rate but not the extent of absorption. The small change in the rate of absorption is not expected to result in a clinically significant effect on steady-state plasma concentrations during once daily dosing of anastrozole tablets. Approximately 90 to 95% of plasma anastrozole steady-state concentrations are attained after 7 daily doses. There is no evidence of time or dose-dependency of anastrozole pharmacokinetic parameters.

Anastrozole pharmacokinetics are independent of age in post-menopausal women.

In boys with pubertal gynecomastia, anastrozole was rapidly absorbed, was widely distributed, and was eliminated slowly with a half-life of approximately 2 days. PK parameters in boys were comparable to those of postmenopausal women. Clearance of anastrozole was lower in girls than in boys and exposure higher. Anastrozole in girls was widely distributed and slowly eliminated, with an estimated half-life of approximately 0.8 days.

Anastrozole is only 40% bound to plasma proteins.

Anastrozole is extensively metabolised by post-menopausal women with less than 10% of the dose excreted in the urine unchanged within 72 hours of dosing. Metabolism of anastrozole occurs by N-dealkylation, hydroxylation and glucuronidation. The metabolites are excreted primarily via the urine. Triazole, a major metabolite in plasma and urine, does not inhibit aromatase.

The apparent oral clearance of anastrozole in volunteers with stable hepatic cirrhosis or renal impairment was in the range observed in healthy volunteers.

## Indication

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Treatment of early breast cancer in hormone receptor positive post-menopausal women.

Adjuvant treatment of early breast cancer in hormone receptor positive postmenopausal women who have received 2 to 3 years of adjuvant tamoxifen.

Reduction in the incidence of contralateral breast cancers in post menopausal women receiving anastrozole as adjuvant treatment for early breast cancer.

Treatment of advanced breast cancer in post-menopausal women.

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## Dosage and Administration

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### *Adults Including The Elderly*

One tablet (1 mg) to be taken orally once a day.

### *Children*

The use of anaccord tablets is not recommended in children, as efficacy has not been established (see ACTIONS and PHARMACOKINETICS).

### *Renal Impairment*

No dose change is recommended.

### *Hepatic Impairment*

No dose change is recommended.

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## Contraindications

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Anaccord Tablets must not be administered during pregnancy or lactation.

Known hypersensitivity to the active substance or to any of the excipients of this product.

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## **Warning & Precautions**

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Anaccord Tablets are not recommended for use in children or in pre-menopausal women as safety and efficacy have not been established in these groups of patients (See ACTIONS AND PHARMACOKINETICS).

Anaccord Tablets have not been investigated in patients with severe hepatic or severe renal impairment. The potential risk/benefit to such patients should be carefully considered before administration of Anaccord Tablets.

As Anastrozole lowers circulating oestrogen levels it may cause a reduction in bone mineral density with a possible consequent increased risk of fracture. This possible increased risk should be managed according to treatment guidelines for managing bone health in postmenopausal women.

### **Use in Pregnancy**

Anaccord Tablets are contraindicated in pregnant women.

### **Use in Lactation**

Anaccord Tablets are contraindicated in breast-feeding women.

### **Effects on Ability to Drive and Use Machines**

Anaccord Tablets are unlikely to impair the ability of patients to drive and operate machinery. However, asthenia and somnolence have been reported with the use of anastrozole and caution should be observed when driving or operating machinery while such symptoms persist.

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## **Adverse Effects**

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Unless specified, the following frequency categories were calculated from the number of adverse events reported in a large phase III study conducted in 9366 postmenopausal women with operable breast cancer treated for 5 years and unless specified, no account was taken of the frequency within the comparative treatment group or whether the investigator considered it to be related to study medication.

Anastrozole Tablets 1 mg

Frequency	System Organ Class	Adverse Reaction
Very common (≥ 10%)	Vascular: General: Musculoskeletal, connective tissue and bone: Nervous system: Gastrointestinal: Skin and subcutaneous tissue:	Hot flushes, mainly mild or moderate in nature. Asthenia, mainly mild or moderate in nature. Joint pain/stiffness, mainly mild or moderate in nature. Headache, mainly mild or moderate in nature. Nausea, mainly mild or moderate in nature. Rash, mainly mild or moderate in nature.
Common (≥ 1% and < 10%)	Skin and subcutaneous tissue: Gastrointestinal: Nervous system: Hepatobiliary disorders: Reproductive system and breast: Metabolism and nutrition:	Hair thinning (Alopecia), mainly mild or moderate in nature. Allergic Reactions Diarrhoea, mainly mild or moderate in nature. Vomiting, mainly mild or moderate in nature. Somnolence, mainly mild or moderate in nature Carpal Tunnel Syndrome* Increases in alkaline phosphatase, alanine aminotransferase and aspartate aminotransferase Vaginal dryness, mainly mild or moderate in nature. Vaginal bleeding, mainly mild or moderate in nature.** Anorexia, mainly mild in nature. Hypercholesterolaemia mainly mild or moderate in nature.
Uncommon (≥ 0.1% and < 1%)	Hepatobiliary disorders: Skin and subcutaneous tissue: Musculoskeletal, connective tissue and bone:	Increases in gamma-GT and bilirubin Hepatitis Urticaria Trigger finger
Rare (≥ 0.01% and < 0.1%)	Skin and subcutaneous tissue:	Erythema multiformae Anaphylactoid reaction
Very rare (< 0.01%)	Skin and subcutaneous tissue:	Stevens-Johnson syndrome Angiodema

\*Events of Carpal Tunnel Syndrome have been reported in patients receiving anastrozole treatment in clinical trials in greater numbers than those receiving treatment with tamoxifen. However, the majority of these events occurred in patients with identifiable risk factors for the development of the condition.

\*\* Vaginal bleeding has been reported commonly, mainly in patients with advanced breast cancer, during the first few weeks after changing from existing hormonal therapy to treatment with Anastrozole. If bleeding persists, further evaluation should be considered.

In a large phase III study conducted in 9366 postmenopausal women with operable breast cancer treated for 5 years, ischaemic cardiovascular events were reported more frequently in patients treated with anastrozole compared to those treated with tamoxifen, although the difference was not statistically significant. The observed difference was mainly due to more reports of angina pectoris and was associated with a sub-group of patients with pre-existing ischaemic heart disease.

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## Interactions

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Antipyrine and cimetidine clinical interaction studies indicate that the co-administration of anastrozole with other medicines is unlikely to result in clinically significant medicine interactions mediated by cytochrome P450.

A review of the clinical trial safety database did not reveal evidence of clinically significant interaction in patients treated with anastrozole who also received other commonly prescribed medicines. There were no clinically significant interactions with biphosphonates (See ACTIONS).

Tamoxifen and/or other therapies containing oestrogen should not be co-administered with anastrozole as they may diminish its pharmacological action.

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## Pharmaceutical Precautions

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### ***Shelf-life and Storage Conditions***

24 months, Store below 25°C.

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## Medicine Classification

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Prescription Medicine

## Package Classification

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Anaccord tablets are presented in a PVC/PVdC aluminium foil blister pack containing 30 tablets

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## Further Information

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### *List of excipients*

- Lactose monohydrate
- Povidone
- Sodium starch glycollate
- Magnesium stearate
- Hypromellose
- Macrogol 300
- Titanium dioxide

### *Pre-clinical Safety Data*

#### **Acute toxicity**

In acute toxicity studies in rodents the median lethal dose of anastrozole was greater than 100 mg/kg by the oral route and greater than 50 mg/kg by the intraperitoneal route. In an oral acute toxicity study in the dog the median lethal dose was greater than 45 mg/kg.

#### **Chronic toxicity**

Multiple dose toxicity studies utilized rats and dogs. No no-effect levels were established for anastrozole in the toxicity studies, but those effects that were observed at the low dose (1 mg/kg/day) and mid doses (dog 3 mg/kg/day; rat 5 mg/kg/day) were related to either the pharmacological or enzyme inducing properties of anastrozole and were unaccompanied by toxic or degenerative changes.

### **Mutagenicity**

Genetic toxicology studies with anastrozole show that it is not a mutagen or a clastogen.

### **Reproductive toxicology**

Oral administration of anastrozole to pregnant rats and rabbits caused no teratogenic effects at doses up to 1.0 and 0.2 mg/kg/day respectively. Those effects that were seen (placental enlargement in rats and pregnancy failure in rabbits) were related to the pharmacology of the compound.

Oral administration of anastrozole to female rats produced a high incidence of infertility at 1 mg/kg/day and increased pre-implantation loss at 0.02 mg/kg/day. These effects were related to the pharmacology of the compound and were completely reversed after a 5-week compound withdrawal period.

The survival of litters born to rats given anastrozole at 0.02 mg/kg/day and above (from day 17 of pregnancy to day 22 post-partum) was compromised. These effects were related to the pharmacological effects of the compound on parturition. There were no adverse effects on behaviour or reproductive performance of the first generation offspring attributable to maternal treatment with anastrozole.

### **Carcinogenicity**

A two year rat oncogenicity study resulted in an increase in incidence of hepatic neoplasms and uterine stromal polyps in females and thyroid adenomas in males at the high dose (25 mg/kg/day) only. These changes occurred at a dose which represents 100-fold greater exposure than occurs at human therapeutic doses, and are considered not to be clinically relevant to the treatment of patients with anastrozole.

A two year mouse oncogenicity study resulted in the induction of benign ovarian tumours and a disturbance in the incidence of lymphoreticular neoplasms (fewer histiocytic sarcomas in females and more deaths as a result of lymphomas). These changes are considered to be mouse-specific effects of aromatase inhibition and not clinically relevant to the treatment of patients with anastrozole.

## **Name and address**

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## **Date of preparation**

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