

# Summary of Product Characteristics

## 1 NAME OF THE MEDICINAL PRODUCT

Alendronic acid/Colecalciferol Rowex 70 mg/2 800 IU tablets

## 2 QUALITATIVE AND QUANTITATIVE COMPOSITION

Each tablet contains 70 mg of alendronic acid (as sodium trihydrate) and 70 microgram (2800 IU) colecalciferol (vitamin D<sub>3</sub>).

### Excipients with known effect:

Each tablet contains 87.1 mg lactose and 13.75 mg sucrose.

For the full list of excipients, see section 6.1.

## 3 PHARMACEUTICAL FORM

Tablet

Oblong-shaped, white to off-white, biconvex tablet, mottled, engraved with '2800' on one side, 12.3 mm in length and 6.5 mm in width.

## 4 CLINICAL PARTICULARS

### 4.1 Therapeutic indications

Alendronic acid/Colecalciferol Rowex is indicated for the treatment of postmenopausal osteoporosis in women at risk of vitamin D insufficiency. It reduces the risk of vertebral and hip fractures.

### 4.2 Posology and method of administration

#### Posology

The recommended dose is one tablet once weekly.

Patients should be instructed that if they miss a dose of Alendronic acid/Colecalciferol Rowex they should take one tablet on the morning after they remember. They should not take two tablets on the same day but should return to taking one tablet once a week, as originally scheduled on their chosen day.

Due to the nature of the disease process in osteoporosis, Alendronic acid/Colecalciferol Rowex is intended for long-term use. The optimal duration of bisphosphonate treatment for osteoporosis has not been established. The need for continued treatment should be re-evaluated periodically based on the benefits and potential risks of Alendronic acid/Colecalciferol Rowex on an individual patient basis, particularly after 5 or more years of use.

Patients should receive supplemental calcium if intake from diet is inadequate (see section 4.4). Additional supplementation with vitamin D should be considered on an individual basis taking into account any vitamin D intake from vitamins and dietary supplements.

The equivalence of intake of 2 800 IU of vitamin D<sub>3</sub> weekly in Alendronic acid/Colecalciferol Rowex to daily dosing of vitamin D 400 IU has not been studied.

#### *Elderly*

In clinical studies there was no age-related difference in the efficacy or safety profiles of alendronate. Therefore no dose adjustment is necessary for the elderly.

#### *Renal impairment*

Alendronic acid/Colecalciferol Rowex is not recommended for patients with renal impairment where creatinine clearance is less than 35 ml/min, due to lack of experience. No dose adjustment is necessary for patients with a creatinine clearance greater than 35 ml/min.

### *Paediatric population*

The safety and efficacy of Alendronic acid/Colecalciferol Rowex in children less than 18 years of age has not been established. This medicinal product should not be used in children less than 18 years of age because no data are available for the alendronic acid/colecalciferol combination. Currently available data for alendronic acid in the paediatric population is described in section 5.1.

### Method of administration

Oral use.

To permit adequate absorption of alendronate:

Alendronic acid/Colecalciferol Rowex must be taken with water only (not mineral water) at least 30 minutes before the first food, beverage, or medicinal product (including antacids, calcium supplements and vitamins) of the day. Other beverages (including mineral water), food and some medicinal products are likely to reduce the absorption of alendronate (see section 4.5 and section 4.8).

The following instructions should be followed exactly in order to minimise the risk of oesophageal irritation and related adverse reactions (see section 4.4):

- Alendronic acid/Colecalciferol Rowex should only be swallowed after getting up for the day with a full glass of water (not less than 200 ml).
- Patients should only swallow Alendronic acid/Colecalciferol Rowex whole. Patients should not crush or chew the tablet or allow the tablet to dissolve in their mouths because of a potential for oropharyngeal ulceration.
- Patients should not lie down for at least 30 minutes after taking Alendronic acid/Colecalciferol Rowex and until after the first food of the day.
- Alendronic acid/Colecalciferol Rowex should not be taken at bedtime or before arising for the day.

### **4.3 Contraindications**

- Hypersensitivity to the active substances or to any of the excipients listed in section 6.1.
- Abnormalities of the oesophagus and other factors which delay oesophageal emptying such as stricture or achalasia.
- Inability to stand or sit upright for at least 30 minutes.
- Hypocalcaemia.

### **4.4 Special warnings and precautions for use**

#### Alendronate

##### *Upper gastrointestinal adverse reactions*

Alendronate can cause local irritation of the upper gastrointestinal mucosa. Because there is a potential for worsening of the underlying disease, caution should be used when alendronate is given to patients with active upper gastrointestinal problems, such as dysphagia, oesophageal disease, gastritis, duodenitis, ulcers, or with a recent history (within the previous year) of major gastrointestinal disease such as peptic ulcer, or active gastrointestinal bleeding, or surgery of the upper gastrointestinal tract other than pyloroplasty (see section 4.3). In patients with known Barrett's oesophagus, prescribers should consider the benefits and potential risks of alendronate on an individual patient basis.

Oesophageal reactions (sometimes severe and requiring hospitalisation), such as oesophagitis, oesophageal ulcers and oesophageal erosions, rarely followed by oesophageal stricture, have been reported in patients receiving alendronate. Physicians should therefore be alert to any signs or symptoms signalling a possible oesophageal reaction and patients should be instructed to discontinue alendronate and seek medical attention if they develop symptoms of oesophageal irritation such as dysphagia, pain on swallowing or retrosternal pain or new or worsening heartburn (see section 4.8).

The risk of severe oesophageal adverse reactions appears to be greater in patients who fail to take alendronate properly and/or who continue to take alendronate after developing symptoms suggestive of oesophageal irritation. It is very important

that the full dosing instructions are provided to, and are understood by the patient (see section 4.2). Patients should be informed that failure to follow these instructions may increase their risk of oesophageal problems.

While no increased risk was observed in extensive clinical trials with alendronate, there have been rare (post-marketing) reports of gastric and duodenal ulcers, some of which were severe and with complications (see section 4.8).

#### *Osteonecrosis of the jaw*

Osteonecrosis of the jaw, generally associated with tooth extraction and/or local infection (including osteomyelitis), has been reported in patients with cancer who are receiving treatment regimens including primarily intravenously administered bisphosphonates. Many of these patients were also receiving chemotherapy and corticosteroids. Osteonecrosis of the jaw has also been reported in patients with osteoporosis receiving oral bisphosphonates.

The following risk factors should be considered when evaluating an individual's risk of developing osteonecrosis of the jaw:

- potency of the bisphosphonate (highest for zoledronic acid), route of administration (see above) and cumulative dose
- cancer, chemotherapy, radiotherapy, corticosteroids, angiogenesis inhibitors, smoking
- a history of dental disease, poor oral hygiene, periodontal disease, invasive dental procedures, and poorly fitting dentures

A dental examination with appropriate preventive dentistry should be considered prior to treatment with oral bisphosphonates in patients with poor dental status.

While on treatment, these patients should avoid invasive dental procedures if possible. For patients who develop osteonecrosis of the jaw while on bisphosphonate therapy, dental surgery may exacerbate the condition. For patients requiring dental procedures, there are no data available to suggest whether discontinuation of bisphosphonate treatment reduces the risk of osteonecrosis of the jaw. Clinical judgement of the treating physician should guide the management plan of each patient based on individual benefit/risk assessment.

During bisphosphonate treatment, all patients should be encouraged to maintain good oral hygiene, receive routine dental check-ups, and report any oral symptoms such as dental mobility, pain, or swelling.

#### *Osteonecrosis of the external auditory canal*

Osteonecrosis of the external auditory canal has been reported with bisphosphonates, mainly in association with long-term therapy. Possible risk factors for osteonecrosis of the external auditory canal include steroid use and chemotherapy and/or local risk factors such as infection or trauma. The possibility of osteonecrosis of the external auditory canal should be considered in patients receiving bisphosphonates who present with ear symptoms such as pain or discharge, or chronic ear infections.

#### *Musculoskeletal pain*

Bone, joint, and/or muscle pain has been reported in patients taking bisphosphonates. In post-marketing experience, these symptoms have rarely been severe and/or incapacitating (see section 4.8). The time to onset of symptoms varied from one day to several months after starting treatment. Most patients had relief of symptoms after stopping treatment. A subset had recurrence of symptoms when rechallenged with the same medicinal product or another bisphosphonate.

#### *Atypical fractures of the femur*

Atypical subtrochanteric and diaphyseal femoral fractures have been reported with bisphosphonate therapy, primarily in patients receiving long-term treatment for osteoporosis. These transverse or short oblique, fractures can occur anywhere along the femur from just below the lesser trochanter to just above the supracondylar flare. These fractures occur after minimal or no trauma and some patients experience thigh or groin pain, often associated with imaging features of stress fractures, weeks to months before presenting with a completed femoral fracture. Fractures are often bilateral; therefore the contralateral femur should be examined in bisphosphonate-treated patients who have sustained a femoral shaft fracture. Poor healing of these fractures has also been reported. Discontinuation of bisphosphonate therapy in patients suspected to have an atypical femur fracture should be considered pending evaluation of the patient, based on an individual benefit risk assessment.

During bisphosphonate treatment patients should be advised to report any thigh, hip or groin pain and any patient presenting with such symptoms should be evaluated for an incomplete femur fracture.

### *Renal impairment*

Alendronic acid/Colecalciferol Rowex is not recommended for patients with renal impairment where creatinine clearance is less than 35 ml/min (see section 4.2).

### *Bone and mineral metabolism*

Causes of osteoporosis other than oestrogen deficiency and ageing should be considered.

Hypocalcaemia must be corrected before initiating therapy with Alendronic acid/Colecalciferol Rowex (see section 4.3). Other disorders affecting mineral metabolism (such as vitamin D deficiency and hypoparathyroidism) should also be effectively treated before starting this medicinal product. The content of vitamin D in Alendronic acid/Colecalciferol Rowex is not suitable for correction of vitamin D deficiency. In patients with these conditions, serum calcium and symptoms of hypocalcaemia should be monitored during therapy with Alendronic acid/Colecalciferol Rowex.

Due to the positive effects of alendronate in increasing bone mineral, decreases in serum calcium and phosphate may occur especially in patients taking glucocorticoids in whom calcium absorption may be decreased. These are usually small and asymptomatic. However, there have been rare reports of symptomatic hypocalcaemia, which have occasionally been severe and often occurred in patients with predisposing conditions (e.g. hypoparathyroidism, vitamin D deficiency and calcium malabsorption) (see section 4.8).

### Colecalciferol

Vitamin D<sub>3</sub> may increase the magnitude of hypercalcaemia and/or hypercalciuria when administered to patients with disease associated with unregulated overproduction of calcitriol (e.g. leukaemia, lymphoma, sarcoidosis). Urine and serum calcium should be monitored in these patients.

Patients with malabsorption may not adequately absorb vitamin D<sub>3</sub>.

### Excipients

This medicinal product contains sucrose. Patients with rare hereditary problems of fructose intolerance, glucose-galactose malabsorption or sucrase-isomaltase insufficiency should not take this medicinal product.

This medicinal product contains lactose. Patients with rare hereditary problems of galactose intolerance, total lactase deficiency or glucose-galactose malabsorption should not take this medicinal product.

This medicinal product contains less than 1 mmol sodium (23 mg) per tablet, that is to say essentially 'sodium-free'.

## **4.5 Interaction with other medicinal products and other forms of interaction**

### Alendronate

If taken at the same time, it is likely that food and beverages (including mineral water), calcium supplements, antacids, and some oral medicinal products will interfere with absorption of alendronate. Therefore, patients must wait at least 30 minutes after taking alendronate before taking any other oral medicinal product (see sections 4.2 and 5.2).

Since Non-Steroidal Anti-Inflammatory Drug (NSAID) use is associated with gastrointestinal irritation, caution should be used during concomitant use with alendronate.

### Colecalciferol

Olestra, mineral oils, orlistat, and bile acid sequestrants (e.g. cholestyramine, colestipol) may impair the absorption of vitamin D. Anticonvulsants, cimetidine and thiazides may increase the catabolism of vitamin D. Additional vitamin D supplements may be considered on an individual basis

## **4.6 Fertility, pregnancy and lactation**

Alendronic acid/Colecalciferol Rowex is only intended for use in postmenopausal women and therefore it should not be used during pregnancy or in breast-feeding women.

### Pregnancy

There are no or limited amount of data from the use of alendronate in pregnant women. Studies in animals have shown reproductive toxicity. Alendronate given during pregnancy in rats caused dystocia related to hypocalcaemia (see section 5.3). Studies in animals have shown hypercalcaemia and reproductive toxicity with high doses of vitamin D (see section 5.3). Alendronic acid/Colecalciferol Rowex should not be used during pregnancy.

### Breast-feeding

It is unknown whether alendronate/metabolites are excreted into human milk. A risk to the newborns/infants cannot be excluded. Colecalciferol and some of its active metabolites pass into breast milk. Alendronic acid/Colecalciferol Rowex should not be used during breast-feeding.

### Fertility

Bisphosphonates are incorporated into the bone matrix, from which they are gradually released over a period of years. The amount of bisphosphonate incorporated into adult bone, and hence, the amount available for release back into the systemic circulation, is directly related to the dose and duration of bisphosphonate use (see section 5.2). There are no data on foetal risk in humans. However, there is a theoretical risk of foetal harm, predominantly skeletal, if a woman becomes pregnant after completing a course of bisphosphonate therapy. The impact of variables such as time between cessation of bisphosphonate therapy to conception, the particular bisphosphonate used, and the route of administration (intravenous versus oral) on the risk has not been studied.

## 4.7 Effects on ability to drive and use machines

Alendronic acid/Colecalciferol Rowex has no or negligible direct influence on the ability to drive and use machines. Patients may experience certain adverse reactions (for example, blurred vision, dizziness and severe bone muscle or joint pain (see section 4.8)) that may influence the ability to drive and use machines.

## 4.8 Undesirable effects

### Summary of the safety profile

The most commonly reported adverse reactions are upper gastrointestinal adverse reactions including abdominal pain, dyspepsia, oesophageal ulcer, dysphagia, abdominal distension and acid regurgitation (> 1 %).

### Tabulated list of adverse reactions

The following adverse reactions have been reported during clinical studies and/or post-marketing use with alendronate.

No additional adverse reactions have been identified for the combination of alendronate and colecalciferol.

Frequencies are defined as: very common ( $\geq 1/10$ ), common ( $\geq 1/100$  to  $< 1/10$ ), uncommon ( $\geq 1/1,000$  to  $< 1/100$ ), rare ( $\geq 1/10,000$  to  $< 1/1,000$ ), very rare ( $< 1/10,000$ ).

System organ class	Frequency	Adverse reactions
<b>Immune system disorders</b>	Rare	hypersensitivity reactions including urticaria and angioedema
<b>Metabolism and nutrition disorders</b>	Rare	symptomatic hypocalcaemia, often in association with predisposing conditions <sup>§</sup>
<b>Nervous system disorders</b>	Common	headache, dizziness <sup>†</sup>
	Uncommon	dysgeusia <sup>†</sup>
<b>Eye disorders</b>	Uncommon	eye inflammation (uveitis, scleritis, or episcleritis)
<b>Ear and labyrinth disorders</b>	Common	vertigo <sup>†</sup>
	Very rare	osteonecrosis of the external auditory canal (bisphosphonate class adverse reaction)
<b>Gastrointestinal disorders</b>	Common	abdominal pain, dyspepsia, constipation, diarrhoea, flatulence, oesophageal ulcer*, dysphagia*, abdominal distension, acid regurgitation
	Uncommon	nausea, vomiting, gastritis, oesophagitis*, oesophageal erosions*, melena <sup>†</sup>
	Rare	oesophageal stricture*, oropharyngeal ulceration*, upper gastrointestinal PUBs (perforation, ulcers, bleeding) <sup>§</sup>
<b>Skin and subcutaneous tissue disorders</b>	Common	alopecia <sup>†</sup> , pruritus <sup>†</sup>
	Uncommon	rash, erythema
	Rare	rash with photosensitivity, severe skin reactions including Stevens-Johnson syndrome and toxic epidermal necrolysis <sup>†</sup>
	Very common	musculoskeletal (bone, muscle or joint) pain which is sometimes

<b>connective tissue disorders</b>		severe <sup>†§</sup>
	Common	joint swelling <sup>†</sup>
	Rare	osteonecrosis of the jaw <sup>†§</sup> ; atypical subtrochanteric and diaphyseal femoral fractures (bisphosphonate class adverse reaction)
<b>General disorders and administration site conditions</b>	Common	asthenia <sup>†</sup> , peripheral oedema <sup>†</sup>
	Uncommon	transient symptoms as in an acute-phase response (myalgia, malaise and rarely, fever), typically in association with initiation of treatment <sup>†</sup>

§See section 4.4

†Frequency in Clinical Trials was similar in the medicinal product and placebo group.

\*See sections 4.2 and 4.4

‡This adverse reaction was identified through post-marketing surveillance. The frequency of rare was estimated based on relevant clinical trials.

### Reporting of suspected adverse reactions

Reporting suspected adverse reactions after authorisation of the medicinal product is important. It allows continued monitoring of the benefit/risk balance of the medicinal product. Healthcare professionals are asked to report any suspected adverse reactions via the national reporting system: HPRA Pharmacovigilance; website: [www.hpra.ie](http://www.hpra.ie)

## 4.9 Overdose

### Alendronate

#### *Symptoms*

Hypocalcaemia, hypophosphataemia and upper gastrointestinal adverse reactions, such as upset stomach, heartburn, oesophagitis, gastritis, or ulcer, may result from oral overdose.

#### *Management*

No specific information is available on the treatment of overdose with alendronate. In case of overdose with Alendronic acid/Colecalciferol Rowex, milk or antacids should be given to bind alendronate. Owing to the risk of oesophageal irritation, vomiting should not be induced and the patient should remain fully upright.

### Colecalciferol

Vitamin D toxicity has not been documented during chronic therapy in generally healthy adults at a dose less than 10,000 IU/day. In a clinical study of healthy adults a 4,000 IU daily dose of vitamin D<sub>3</sub> for up to five months was not associated with hypercalciuria or hypercalcaemia.

## 5 PHARMACOLOGICAL PROPERTIES

### 5.1 Pharmacodynamic properties

Pharmacotherapeutic group: Drugs for treatment of bone diseases, Bisphosphonates, combinations, ATC code: M05BB03

#### Mechanism of action

##### *Alendronate*

Alendronate sodium is a bisphosphonate that inhibits osteoclastic bone resorption with no direct effect on bone formation. Preclinical studies have shown preferential localisation of alendronate to sites of active resorption. Activity of osteoclasts is inhibited, but recruitment or attachment of osteoclasts is not affected. The bone formed during treatment with alendronate is of normal quality.

##### *Colecalciferol (vitamin D<sub>3</sub>)*

Vitamin D<sub>3</sub> is produced in the skin by conversion of 7-dehydrocholesterol to vitamin D<sub>3</sub> by ultraviolet light. In the absence of adequate sunlight exposure, vitamin D<sub>3</sub> is an essential dietary nutrient. Vitamin D<sub>3</sub> is converted to 25-hydroxyvitamin D<sub>3</sub> in the liver, and stored until needed. Conversion to the active calcium-mobilising hormone 1,25-dihydroxyvitamin D<sub>3</sub> (calcitriol) in the kidney is tightly regulated. The principal action of 1,25-dihydroxyvitamin D<sub>3</sub> is to increase intestinal absorption of both calcium and phosphate as well as regulate serum calcium, renal calcium and phosphate excretion, bone formation and bone resorption.

Vitamin D<sub>3</sub> is required for normal bone formation. Vitamin D insufficiency develops when both sunlight exposure and dietary intake are inadequate. Insufficiency is associated with negative calcium balance, bone loss, and increased risk of skeletal fracture. In severe cases, deficiency results in secondary hyperparathyroidism, hypophosphataemia, proximal muscle weakness and osteomalacia, further increasing the risk of falls and fractures in osteoporotic individuals. Supplemental vitamin D reduces these risks and their consequences.

Osteoporosis is defined as bone mineral density (BMD) of the spine or hip 2.5 standard deviations (SD) below the mean value of a normal young population or as a previous fragility fracture, irrespective of BMD.

### Clinical efficacy and safety

#### *Alendronate/colecalciferol studies*

The effect of the lower dose of alendronate 70 mg/vitamin D<sub>3</sub> 2 800 IU on vitamin D status was demonstrated in a 15-week, multinational study that enrolled 682 osteoporotic post-menopausal women (serum 25-hydroxyvitamin D at baseline: mean, 56 nmol/l [22.3 ng/ml]; range, 22.5-225 nmol/l [9-90 ng/ml]). Patients received the lower strength (70 mg/2 800 IU) of alendronate/vitamin D<sub>3</sub> (n=350) or alendronate 70 mg (n=332) once a week; additional vitamin D supplements were prohibited. After 15 weeks of treatment, the mean serum 25-hydroxyvitamin D levels were significantly higher (26 %) in the alendronate/vitamin D<sub>3</sub> (70 mg/2 800 IU) group (56 nmol/l [23 ng/ml]) than in the alendronate-only group (46 nmol/l [18.2 ng/ml]). The percentage of patients with vitamin D insufficiency (serum 25-hydroxyvitamin D < 37.5 nmol/l [< 15 ng/ml]) was significantly reduced by 62.5 % with alendronate/vitamin D<sub>3</sub> (70 mg/2 800 IU) vs. alendronate-only (12 % vs. 32 %, respectively), through week 15. The percentage of patients with vitamin D deficiency (serum 25-hydroxyvitamin D < 22.5 nmol/l [< 9 ng/ml]) was significantly reduced by 92 % with alendronate/vitamin D<sub>3</sub> (70 mg/2 800 IU) vs. alendronate-only (1 % vs 13 %, respectively). In this study, mean 25-hydroxyvitamin D levels in patients with vitamin D insufficiency at baseline (25-hydroxyvitamin D, 22.5 to 37.5 nmol/l [9 to < 15 ng/ml]) increased from 30 nmol/l (12.1 ng/ml) to 40 nmol/l (15.9 ng/ml) at week 15 in the alendronate/vitamin D<sub>3</sub> (70 mg/2 800 IU) group (n=75) and decreased from 30 nmol/l (12.0 ng/ml) at baseline to 26 nmol/l (10.4 ng/ml) at week 15 in the alendronate-only group (n=70). There were no differences in mean serum calcium, phosphate, or 24-hour urine calcium between treatment groups.

The effect of the lower dose of alendronate 70 mg/vitamin D<sub>3</sub> 2 800 IU plus an additional 2 800 IU vitamin D<sub>3</sub> for a total of 5 600 IU (the amount of vitamin D<sub>3</sub> in the higher dose of alendronate/vitamin D<sub>3</sub>) once weekly was demonstrated in a 24-week, extension study that enrolled 619 osteoporotic post-menopausal women. Patients in the vitamin D<sub>3</sub> 2 800 group received alendronate/vitamin D<sub>3</sub> (70 mg/2 800 IU) (n=299) and patients in the vitamin D<sub>3</sub> 5 600 group received alendronate/vitamin D<sub>3</sub> (70 mg/2 800 IU) plus an additional 2 800 IU vitamin D<sub>3</sub> (n=309) once a week; additional vitamin D supplements were allowed. After 24-weeks of treatment, the mean serum 25-hydroxyvitamin D levels were significantly higher in the vitamin D<sub>3</sub> 5 600 group (69 nmol/l [27.6 ng/ml]) than in the vitamin D<sub>3</sub> 2 800 group (64 nmol/l [25.5 ng/ml]). The percentage of patients with vitamin D insufficiency was 5.4 % in the vitamin D<sub>3</sub> 2 800 group vs. 3.2 % in the vitamin D<sub>3</sub> 5 600 group through the 24-week extension. The percentage of patients with vitamin D deficiency was 0.3 % in the Vitamin D<sub>3</sub> 2 800 group vs. zero in the Vitamin D<sub>3</sub> 5 600 group. There were no differences in mean serum calcium, phosphate, or 24-hour urine calcium between treatment groups. The percentage of patients with hypercalciuria at the end of the 24-week extension was not statistically different between treatment groups.

#### *Alendronate studies*

The therapeutic equivalence of alendronate once weekly 70 mg (n=519) and alendronate 10 mg daily (n=370) was demonstrated in a one-year multicentre study of post-menopausal women with osteoporosis. The mean increases from baseline in lumbar spine BMD at one year were 5.1 % (95 % CI: 4.8, 5.4 %) in the 70 mg once-weekly group and 5.4 % (95 % CI: 5.0, 5.8 %) in the 10 mg daily group. The mean BMD increases were 2.3 % and 2.9 % at the femoral neck and 2.9 % and 3.1 % at the total hip in the 70 mg once weekly and 10 mg daily groups, respectively. The two treatment groups were also similar with regard to BMD increases at other skeletal sites.

The effects of alendronate on bone mass and fracture incidence in post-menopausal women were examined in two initial efficacy studies of identical design (n=994) as well as in the Fracture Intervention Trial (FIT: n=6,459).

In the initial efficacy studies, the mean BMD increases with alendronate 10 mg/day relative to placebo at three years were 8.8%, 5.9 % and 7.8 % at the spine, femoral neck and trochanter, respectively. Total body BMD also increased significantly. There was a 48 % reduction (alendronate 3.2 % vs placebo 6.2 %) in the proportion of patients treated with alendronate experiencing one or more vertebral fractures relative to those treated with placebo. In the two-year extension of these studies BMD at the spine and trochanter continued to increase and BMD at the femoral neck and total body were maintained.

FIT consisted of two placebo-controlled studies using alendronate daily (5 mg daily for two years and 10 mg daily for either one or two additional years):

- FIT 1: A three-year study of 2,027 patients who had at least one baseline vertebral (compression) fracture. In this study alendronate daily reduced the incidence of  $\geq 1$  new vertebral fracture by 47 % (alendronate 7.9 % vs. placebo 15.0 %). In addition, a statistically significant reduction was found in the incidence of hip fractures (1.1% vs. 2.2 %, a reduction of 51 %).
- FIT 2: A four year study of 4,432 patients with low bone mass but without a baseline vertebral fracture. In this study, a significant difference was observed in the analysis of the subgroup of osteoporotic women (37 % of the global population who correspond with the above definition of osteoporosis) in the incidence of hip fractures (alendronate 1.0 % vs. placebo 2.2 %, a reduction of 56 %) and in the incidence of  $\geq 1$  vertebral fracture (2.9 % vs 5.8 %, a reduction of 50 %).

#### *Laboratory test findings*

In clinical studies, asymptomatic, mild and transient decreases in serum calcium and phosphate were observed in approximately 18 % and 10 %, respectively, of patients taking alendronate 10 mg/day versus approximately 12 % and 3 % of those taking placebo. However, the incidences of decreases in serum calcium to  $< 8.0$  mg/dl (2.0 mmol/l) and serum phosphate to  $\leq 2.0$  mg/dl (0.65 mmol/l) were similar in both treatment groups.

#### Paediatric population

Alendronate sodium has been studied in a small number of patients with osteogenesis imperfecta under the age of 18 years. Results are insufficient to support the use of alendronate sodium in paediatric patients with osteogenesis imperfecta.

## **5.2 Pharmacokinetic properties**

### Alendronate

#### Absorption

Relative to an intravenous reference dose, the oral mean bioavailability of alendronate in women was 0.64 % for doses ranging from 5 to 70 mg when administered after an overnight fast and two hours before a standardised breakfast. Bioavailability was decreased similarly to an estimated 0.46 % and 0.39 % when alendronate was administered one hour or half an hour before a standardised breakfast. In osteoporosis studies, alendronate was effective when administered at least 30 minutes before the first food or beverage of the day.

The alendronate component in the Alendronic acid/Colecalciferol Rowex (70 mg/2 800 IU) combination tablet and the alendronate/vitamin D<sub>3</sub> (70 mg/5 600 IU) combination tablet is bioequivalent to the alendronate 70 mg tablet.

Bioavailability was negligible whether alendronate was administered with, or up to two hours after, a standardised breakfast. Concomitant administration of alendronate with coffee or orange juice reduced bioavailability by approximately 60 %.

In healthy subjects, oral prednisone (20 mg three times daily for five days) did not produce a clinically meaningful change in oral bioavailability of alendronate (a mean increase ranging from 20 % to 44 %).

#### Distribution

Studies in rats show that alendronate transiently distributes to soft tissues following 1 mg/kg intravenous administration but is then rapidly redistributed to bone or excreted in the urine. The mean steady-state volume of distribution, exclusive of bone, is at least 28 litres in humans. Concentrations of alendronate in plasma following therapeutic oral doses are too low for analytical detection ( $< 5$  ng/ml). Protein binding in human plasma is approximately 78 %.

#### Biotransformation

There is no evidence that alendronate is metabolised in animals or humans.

#### Elimination

Following a single intravenous dose of [<sup>14</sup>C] alendronate, approximately 50 % of the radioactivity was excreted in the urine within 72 hours and little or no radioactivity was recovered in the faeces. Following a single 10 mg intravenous dose, the renal clearance of alendronate was 71 ml/min, and systemic clearance did not exceed 200 ml/min. Plasma concentrations fell by more than 95 % within six hours following intravenous administration. The terminal half-life in humans is estimated to exceed ten years, reflecting release of alendronate from the skeleton. Alendronate is not excreted

through the acidic or basic transport systems of the kidney in rats, and thus it is not anticipated to interfere with the excretion of other medicinal products by those systems in humans.

Colecalciferol.

#### Absorption

In healthy adult subjects (males and females), following administration of Alendronic acid/Colecalciferol Rowex 70 mg/2 800 IU tablets after an overnight fast and two hours before a meal, the mean area under the serum-concentration-time curve (AUC<sub>0-120 hrs</sub>) for vitamin D<sub>3</sub> (unadjusted for endogenous vitamin D<sub>3</sub> levels) was 296.4 ng-hr/ml. The mean maximal serum concentration (C<sub>max</sub>) of vitamin D<sub>3</sub> was 5.9 ng/ml, and the median time to maximal serum concentration (T<sub>max</sub>) was 12 hours. The bioavailability of the 2 800 IU vitamin D<sub>3</sub> in alendronate/vitamin D<sub>3</sub> is similar to 2 800 IU vitamin D<sub>3</sub> administered alone.

In healthy adult subjects (males and females), following administration of alendronate/vitamin D<sub>3</sub> 70 mg/5 600 IU after an overnight fast and two hours before a meal, the mean area under the serum- concentration- time curve (AUC<sub>0-80 hrs</sub>) for vitamin D<sub>3</sub> (unadjusted for endogenous vitamin D<sub>3</sub> levels) was 490.2 ng-hr/ml. The mean maximal serum concentration (C<sub>max</sub>) of vitamin D<sub>3</sub> was 12.2 ng/ml and the median time to maximal serum concentration (T<sub>max</sub>) was 10.6 hours. The bioavailability of the 5 600 IU vitamin D<sub>3</sub> in alendronate/vitamin D<sub>3</sub> is similar to 5 600 IU vitamin D<sub>3</sub> administered alone.

#### Distribution

Following absorption, vitamin D<sub>3</sub> enters the blood as part of chylomicrons. Vitamin D<sub>3</sub> is rapidly distributed mostly to the liver where it undergoes metabolism to 25-hydroxyvitamin D<sub>3</sub>, the major storage form. Lesser amounts are distributed to adipose and muscle tissue and stored as vitamin D<sub>3</sub> at these sites for later release into the circulation. Circulating vitamin D<sub>3</sub> is bound to vitamin D-binding protein.

#### Biotransformation

Vitamin D<sub>3</sub> is rapidly metabolised by hydroxylation in the liver to 25-hydroxyvitamin D<sub>3</sub>, and subsequently metabolised in the kidney to 1,25-dihydroxyvitamin D<sub>3</sub>, which represents the biologically active form. Further hydroxylation occurs prior to elimination. A small percentage of vitamin D<sub>3</sub> undergoes glucuronidation prior to elimination.

#### Elimination

When radioactive vitamin D<sub>3</sub> was administered to healthy subjects, the mean urinary excretion of radioactivity after 48 hours was 2.4 %, and the mean faecal excretion of radioactivity after 4 days was 4.9 %. In both cases, the excreted radioactivity was almost exclusively as metabolites of the parent. The mean half-life of vitamin D<sub>3</sub> in the serum following an oral dose of alendronate acid/vitamin D<sub>3</sub> (70 mg/2 800 IU) is approximately 24 hours.

#### Renal impairment

Preclinical studies show that alendronate that is not deposited in bone is rapidly excreted in the urine. No evidence of saturation of bone uptake was found after chronic dosing with cumulative intravenous doses up to 35 mg/kg in animals. Although no clinical information is available, it is likely that, as in animals, elimination of alendronate via the kidney will be reduced in patients with impaired renal function. Therefore, somewhat greater accumulation of alendronate in bone might be expected in patients with impaired renal function (see section 4.2).

### **5.3 Preclinical safety data**

Non-clinical studies with the combination of alendronate and colecalciferol have not been conducted.

#### Alendronate

Non-clinical data reveal no special hazard for humans based on conventional studies of safety pharmacology, repeated dose toxicity, genotoxicity and carcinogenic potential. Studies in rats have shown that treatment with alendronate during pregnancy was associated with dystocia in dams during parturition which was related to hypocalcaemia. In studies, rats given high doses showed an increased incidence of incomplete foetal ossification. The relevance to humans is unknown.

#### Colecalciferol

At doses far higher than the human therapeutic range, reproductive toxicity has been observed in animal studies.

## **6 PHARMACEUTICAL PARTICULARS**

### **6.1 List of excipients**

Lactose  
Microcrystalline cellulose (E460)  
Croscarmellose sodium (E468)  
Magnesium stearate (E572)  
Sunflower oil, refined  
Butylhydroxytoluene (E321)  
Gelatin  
Sucrose  
Maize starch  
Aluminium magnesium silicate

### **6.2 Incompatibilities**

Not applicable.

### **6.3 Shelf life**

2 years.

### **6.4 Special precautions for storage**

This medicinal product does not require any special temperature storage conditions. Store in the original package in order to protect from moisture and light.

### **6.5 Nature and contents of container**

oPA/Alu/PVC//Alu blister  
Pack sizes: cartons containing 2, 4, 6 or 12 tablets.

Not all pack sizes may be marketed.

### **6.6 Special precautions for disposal**

Any unused medicinal product or waste material should be disposed of in accordance with local requirements.

## **7 MARKETING AUTHORISATION HOLDER**

Rowex Ltd  
Newtown  
Bantry  
Co. Cork  
Ireland

## **8 MARKETING AUTHORISATION NUMBER**

PA0711/319/001

## **9 DATE OF FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION**

Date of Authorisation: 31<sup>st</sup> March 2023

## **10 DATE OF REVISION OF THE TEXT**