Proton pump inhibitors in childhood cystic fibrosis

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**PROTON PUMP INHIBITORS IN
CHILDHOOD CYSTIC FIBROSIS**

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LIST OF ABBREVIATIONS

BHR	bronchial hyper responsiveness	PL	phospholipids
BL	bronchial lability	PUFA	poly unsaturated fatty acid
BMC	bone mineral content	PPI	proton pump inhibitor
BMD	bone mineral density	(s)RAW	(specific) airway resistance
BMI	body mass index	RBC	red blood cell
CF	cystic fibrosis	RDA	recommended daily allowance
CFTR	CF transmembrane regulator (protein)	REE	resting energy expenditure
COPD	chronic obstructive pulmonary disease	RV	residual volume
DEXA	dual energy x-ray absorptio metry	SDS	standard deviation score
ERV	expiratory reserve volume	SFA	saturated fatty acid
ECFS	European CF Society	TBF	total body fat
ENT	ear – nose – throat	TLC	total lung capacity
ESCF	epidemiologic study of CF		
(E)FA	(essential) fatty acid		
FFM	fat free mass		
FM	fat mass		
FEV ₁	forced expiratory volume in one second		
FVC	forced vital capacity		
(s)GAW	(specific) airway conductance		
GER(D)	gastro esophageal reflux disease		
GI	gastro intestinal		
(t)Hcy	(total) homocysteine		
ITGV	intra thoracic gas volume		
IP	intestinal permeability		
II	intestinal inflammation		
Lcp	long chain poly unsaturated FA		
L/M/Rh	lactulose – mannitol – L-rhamnose		
NO(x)	nitric oxide (oxidation products)		
NS	not significant		
MIF _{50%VCin}	maximal inspiratory flow halfway VCin		
MUFA	mono unsaturated fatty acid		
PEF	peak expiratory flow		
PImax	maximal inspiratory mouth pressure		
PEmax	maximal expiratory mouth pressure		
PERT	pancreatic enzyme replacement therapy		

CHAPTER ONE

Rationale for the use of Proton Pump Inhibitors in CF patients

Han Hendriks and Philippe Forget

1.1. GENERAL INTRODUCTION

Cystic Fibrosis (CF) is one of the most common lethal genetic diseases in Caucasians, affecting approximately 1 in 3,400 live births [1, 2]. A similar incidence has been reported in the Netherlands [3]. Although this autosomal recessive inherited disorder affects only one gene, a heterogenous phenotypic presentation of CF is seen. This can be explained by the many different mutations of this single gene, and by the presence of the so-called modifier genes interacting with the affected gene. The genetic defect causing the primary disease affects the cystic fibrosis transmembrane conductance regulator protein (CFTR), leading to a defect in the chloride and bicarbonate transport regulating channel at the luminal side of exocrine cells [4].

Since the first description of CF as a separate disease entity by Fanconi in 1936 and Anderson in 1938, many other researchers have contributed to the present knowledge of the pathophysiology of this disease. Besides the name Cystic Fibrosis, the term Mucoviscoidosis is also commonly used. This term refers to one of the most striking abnormalities present in CF patients, namely the insufficient clearance of increased amounts of viscous mucus from the lungs. The designation "Mucoviscoidosis" has been translated in many languages, for example in Dutch "taai-slijm-ziekte", but many other terms are being used worldwide.

It was not until 1989 that the gene, which encodes for the cAMP dependent chloride channel, the CFTR, was first identified [5-8]. This discovery led to a new direction, and an explosive growth, in CF research improving our knowledge of the basal pathophysiology. In this review we will try to describe in detail some new insights in the pathophysiology of CF as well as the rationale behind the use of proton pump inhibitors (PPI) in CF patients.

1.2. PATHOPHYSIOLOGY OF CYSTIC FIBROSIS

Most patients with CF are born with (>65%) or acquire (>90%) insufficiency of exocrine pancreatic function soon after birth. Nearly all have at that time a sweat chloride concentration of more than 60 mMol/L and only 1 out of thousand typical CF patients will have a lower value. At presentation lung involvement is reported to be less than 40%. However, pulmonary problems causes most of the morbidity and nearly all mortality in CF patients. An abnormal sweat test together with pulmonary and pancreatic disease forms the classical triad of findings confirming the clinical diagnosis of CF [9]. All these patients have mutations in both copies of the CFTR gene on the long arm of chromosome 7, but not everyone with two mutant alleles will manifest the full CF syndrome. Meanwhile over 1000 different mutations have been identified, of which $\Delta F508$ is the most common (more than 70% of the mutations in Caucasians from Northern and Western European origin). These different mutations, coding for the same single protein of about 250 kb of genomic DNA, containing 27 exons, are classified in 5 different classes. While for pancreatic insufficiency there is a fairly good correlation between phenotype and genotype, this is not the case for the pul-

monary involvement.[10] This has led to the concept of so-called modifier genes present on other locations and in other chromosomes. This may result in totally different phenotypes of CF, within the same genotype of one family. This is perhaps the most important reason why a given set of CF mutations cannot reliably predict the prognosis. This makes it difficult not only to describe one common pathophysiologic process in all CF patients, but also to compare CF patients with each other.

At least two basic functional abnormalities are attributable to a dysfunctional CFTR. Not only does this involve a defective, energy-dependent, anion channel in apical epithelial membranes, but it also affects other membrane proteins, regulating other ion channels allowing ions to flow either in or out of cells. Therefore CFTR is crucial in both absorption and secretion of ions, especially in wet epithelia [4]. As a result of this basic mechanism two different, not necessarily conflicting, pathophysiologic hypotheses have been developed. First, the so-called low volume hypothesis suggests that the decreased water content of the secretions provides a sticky, viscous product, easily blocking the excretory ducts. This results in an impaired ciliary functioning, with decreased or absent mucociliary clearance, allowing a state of permanent bacterial colonisation and infection of the airways, inducing inflammation [11]. Second, the high salt hypothesis describes the condition where high concentrations of chloride and sodium, as seen in CF, impair normal functioning of small peptides floating in the epithelial lining fluid. Some of these peptides play an important role in host defence against bacteria and, if defective, will give rise to the growth of bacteria, in particular *Pseudomonas Aeruginosa* and *Burkholderia Cepacia*, bacteria which are commonly cultured in sputum of CF patients [12].

As mucosal exocrine glands from various organs are differently affected, this can explain why loss of CFTR function will have differing consequences depending upon the cell type involved. Furthermore, findings, for instance in the vas deferens in man and mice, show that in different species the same gland with the same defective CFTR may be affected differently [13, 14].

1.3. THE LINK BETWEEN NUTRITIONAL CONDITION AND PROGNOSIS IN CF

In chronic obstructive pulmonary disease (COPD) a deterioration of the nutritional condition, as measured by decreased body mass index (BMI)[15], fat free mass (FFM) is correlated with a poor prognosis, independently of airflow obstruction and clinical symptoms [16-20]. Because of this relationship it may be that the decrease in body composition precedes the decline of lung function [21]. Despite the clinical similarity between COPD and CF, regarding pulmonary involvement and its relation with body composition, there are remarkable differences between those diseases. It has been shown in COPD that an abnormal protein metabolism, as reflected by lower plasma branched-chain amino acid concentrations and different muscle amino acid patterns, is responsible for the decreased FFM [16, 22, 23]. As mentioned in the previous section, in CF there seems to be a depletion in fat mass (FM) rather than an abnormal

FFM as found in COPD. In both disorders, however, FFM is decreased, resulting in problems of endurance of skeletal muscles and in particular of respiratory muscles, where inspiratory muscles are more affected than expiratory muscles [24, 25]. Besides a direct impairment of the respiratory muscles, an increase of inflammatory markers has been found to be associated with decreasing FFM in COPD [26]. Serum levels of leptin, an important regulating peptide for energy homeostasis which increases body weight, has been shown to correlate with FFM in COPD [27]. Whether or not such a relationship also exists in CF is unknown. Several researchers have studied the relationship between nutritional condition, lung function and survival in CF patients [28, 29]. In some CF subgroups, such as pre-lung-transplant patients and pregnant patients, this relationship has been established [30-32]. Although strongly suspected, a direct relation between nutritional condition, lung function and survival in the whole CF population has, until now, not been unequivocally proven. Even the large epidemiologic study of CF (ESCF) [33, 34] was unable to demonstrate a significant correlation between lung disease and nutritional status. Only in adolescent and young adult females, due to a survivor effect, could some relationship be shown. The cross-sectional design of the ESCF was held responsible for the lack of correlation between nutritional status and lung disease. Studies supporting a direct relation between lung function and nutritional status in CF are sparse [35, 36]. There is a strong need for a large longitudinal, long-term, placebo-controlled study to evaluate the effect of therapies improving nutritional status on lung function and survival. Such a study, however, is difficult to perform; it should be multicentric, and consequently would be very expensive.

1.4. FACTORS AFFECTING NUTRITIONAL CONDITION OF CF PATIENTS

Low energy intake, high fecal losses and increased energy expenditure can all contribute to a poor nutritional condition in CF.

INTAKE

Studies conducted in the 1980s have shown that energy and fat enriched diets are mandatory to improve survival in CF patients [37-39]. In CF infants with pancreatic insufficiency, there is no need for (semi-) elemental predigested formula feeding, because absorption of these is not better than that of ordinary formulas, administered with enzyme replacement [40]. Because of ongoing stool losses, it is recommended to provide a high fat diet (35-40% total energy), which, in most patients, results in normal growth for height and weight until their lung disease deteriorates significantly. Patients who develop a negative energy balance seldom if ever respond to such a dietary therapy and will need supplemental, nocturnal, gastrostomy feeds, to improve nutritional status [41]. One should be aware that CF children often cannot consume the required intake of 120% of recommended dietary allowances (RDA) for both energy and fat. Energy intakes are often insufficient to meet requirements for normal growth [42]. The Cochrane library has collected evidence of the benefit of oral

protein and calorie supplements to improve the nutritional status of children with chronic diseases. Only three trials assessed the effects of such food supplements in children with CF and apparently it was not possible to draw any firm conclusions concerning the benefit of these therapies [43, 44]. We therefore suggest that the improvement in the nutritional status reported in the 1980s was mainly due to improvement of malabsorption by introducing (enteric-coated) pancreatic enzymes and not so much the result of increased dietary fat intake.

MALABSORPTION

While the main cause of malabsorption in CF is a deficiency of pancreatic enzymes, bicarbonate deficiency, abnormalities of bile salts, impaired mucosal transport and motility, as well as anatomical structural changes are contributory factors too. Appropriate pancreatic enzyme replacement therapy will achieve satisfactory absorption in many patients. Therefore it is important to identify both malabsorption and evidence of pancreatic insufficiency in all CF patients. All those who have evidence of fat malabsorption are considered pancreatic insufficient and require enzyme replacement therapy. [45]. As long ago as the early 1960s exogenous pancreatic enzymes were used in an attempt to decrease steatorrhea. Relatively crude extracts of porcine pancreas with low enzyme potency were used initially [46]. Later several different enteric-coated enzymes were introduced [47]. Pancreatic enzyme replacement therapy (PERT) is a major factor for achieving normal growth and nutritional status in CF patients with pancreatic insufficiency. Combining PERT with supplemental energy intake further improves the nutritional condition. This approach should also include patient education to improve compliance with treatment. Patients and their families will have a better understanding of the relationship between PERT and nutritional status if appropriate guidelines are given and correctly applied [48].

ENERGY EXPENDITURE (EE)

CFTR mRNA is expressed in the human anterior hypothalamus, an area involved in regulation of appetite, resting energy expenditure (REE) and sexual differentiation [49]. It has been suggested that in CF there is an energy-wasting lesion that will increase total EE (TEE) and hence energy requirement. Recent data, with the double labelled water technique, collected from babies with CF strongly suggest that this is not the case. It is probable that previous data were confounded by heterogeneity of disease status [50]. In CF children with mild to moderate lung disease, an acute pulmonary exacerbation has not been found to be associated with an increase of REE [51, 52]. Therefore it has been concluded that no direct relationship between the increase of basic metabolic rate, as assessed by REE in CF, and nutritional status or pulmonary disease can be demonstrated [53].

1.5. PANCREATIC INSUFFICIENCY AND ENZYME SUPPLETION IN CF

As already mentioned, one of the most serious consequences of an impaired CFTR function is pancreatic insufficiency, causing maldigestion and in particular fat malabsorption. This pancreatic insufficiency is only partially corrected by exogenous pancreatic enzymes [54]. Increasing the dose of these enzymes might result in serious complications like fibrosing colonopathy, where pancreatic enzymes if still active in the colon are held responsible for scarring of the colon tissue [55].

In most patients with CF the pancreas is gradually destroyed and replaced by fibrous cysts. This process already starts in fetal life [56]. Normally, exocrine pancreatic gland cells secrete digestive enzymes which mix with a bicarbonate-rich fluid secreted by duct cells, flowing into the duodenum. The production of the ductal juice is probably first reduced, and not the enzyme production itself, causing the ducts to be blocked by enzyme precipitates leading to the destruction of excretory ducts [57, 58]. The exocrine pancreas is different from other exocrine glands in that its acini entirely lack myoepithelial cells supporting the ducts. Hence they are sensitive to damage by increases in pressure that occur when the duct is blocked. Mutations that spare some CFTR function (estimated to be more than 5% of normal) leave enough secretion to maintain patency of the ductular system and thus avoid destruction of the pancreas, preventing pancreatic insufficiency. However, the vast majority of CF patients will be pancreatic insufficient, from the first year of life, resulting in a variable degree of growth failure. While the cause of malnutrition seems clear, the degree of impaired digestion and absorption of nutrients will determine the severity of it. Normally the pancreas produces several types of enzymes: lipase, necessary for the absorption of lipids, proteases, for the cleavage of proteins, and amylase, for the hydrolysis of large carbohydrate molecules. These enzymes are secreted, together with bicarbonate, into the intestine. Although secretion of all these types of enzymes by the pancreas will be equally reduced, secretion of the gastrointestinal (GI) tract will compensate for deficiencies in amylases and proteases, in contrast to lipase. Deficiency of lipase will result in decreased fat absorption and steatorrhea, resulting in failure to grow. The complete pathophysiology of fat malabsorption is much more complicated. This is apparent from the fact that the addition of exogenous pancreatic enzymes alone does improve but does not normalize malabsorption completely. Increasing the dose of enzymes does not result in an equivalent increase of nutritional absorption, and will lead to a plateau of maximal effect, most often beneath the level of normalization. Further addition of enzymes does not give any further improvement, while it can induce complications such as fibrosing colonopathy [59].

The lack of normalization of steatorrhea despite sufficient suppletion of exogenous pancreatic enzymes has been related to several factors:

1. Mixing between enzymes from capsules and food is never as optimal as in vivo.
2. The orally supplemented enzymes need to be supplied with an enteric-coating in order to resist destruction by gastric acid. If released from these capsules in an acid milieu, they will be denatured, which makes them inactive. Lipase, for instance, is irreversibly destroyed at a pH of 4.0 or lower [60].

3. Although there are some small differences between preparations, the enteric-coating capsules dissolves at a pH above 5.5. [61] Normally this will take place in the proximal duodenum, where as a result of production of bicarbonate by the pancreatic ducts, gastric acid is buffered. As already mentioned, it is this lack of buffering capacity of the pancreas, which occurs in a very early phase of the disease. Consequently, the pH of the duodenal contents will be lower than 5.5 for a variable but longer distance in the GI tract compared to healthy persons [62]. Dissolution of the coating will therefore be delayed, while the transit of the contents continues. Therefore a valuable mucosal absorption area will be left unutilized, especially for fat absorption, as this takes place in the duodenum and proximal jejunum [63].
4. A well-known finding in CF is an important increase of intestinal permeability (IP). [64]. It might be that the acid intestinal contents affect the intestinal mucosa, increasing IP, which might provoke several undesired effects such as allergic reactions or increased intestinal inflammation, worsening malabsorption and abdominal complaints.
5. Ionised bile salts, in a sufficient concentration, are necessary to form micelles. In CF the output of bile salts is decreased because of precipitation in the bile ducts, and with the steatorrhea secreted bile salts are lost in the stools. If the concentration of these bile salts is lower than required to form micelles, the critical micelles concentration (CMC), this will lead to reduced transport of the lipolytic products to the microvilli for absorption. An increasing number of papers present evidence that supplementation of bile salts (ursodeoxycholic acid) to CF patients will not only diminish the degree of cirrhosis but will also improve fat malabsorption [65, 66].

Reviewing the literature (1966-1994) concerning pH, motility/transit, and permeability of the GI tract in CF, Gregory [67] reported that most studies were performed with very small numbers of patients. But even when considered the published data together, they do not confirm some generally expressed views on these topics. The only consistent observations were:

- ◆ A high incidence of gastroesophageal reflux in CF
- ◆ Pre- and postprandial duodenal pH 1-2 units lower in CF patients compared to healthy controls
- ◆ Small intestinal paracellular permeability 4-10 times greater than normal in CF.

A decrease in duodenal pH in CF has been evaluated by Barraclough and Taylor [54]. By assessing gastric and proximal duodenal pH, during 24-h ambulatory pH measurements, they showed that children with CF had a normal fasting gastric and duodenal pH, while their postprandial duodenal pH was markedly decreased compared to that of controls. This became even more pronounced with successive meals. The total time that duodenal pH was <5 varied from 15-90% of the recordings (mean 57%). Duodenal pH returned to normal levels only overnight, in the fasting state. The pancreas has an enormous reserve capacity, so significant malabsorption most often signals severe reduction of exocrine function. Complete duodenal hydrolysis of fat requires a series of complex interdependent physicochemical events involving pancreatic lipase, colipase,

phospholipase A2 and bile salts in an environment where the pH must be close to neutrality [68]. Digestion of lipids can be described by a chain of events: emulsification, lipolysis, solubilization, uptake with re-esterification in the enterocyte, and transportation into the circulation via the lymph. Defects at any of these stages can cause steatorrhea.

The low pH may also contribute to the development of fibrosing colonopathy. High dose pancreatic enzyme preparations might fail to dissolve because of the acidity in the small intestinal lumen, leaving unutilized proteolytic enzymes enter the colon. Adherence to the colonic wall might lead to destruction, with scarring and strictures of the colonic wall.

1.6. THERAPEUTICAL APPROACHES TO OPTIMIZE THE EFFECT OF EXOGENOUS PANCREATIC ENZYMES

ACID LIPASES

Acid duodenal contents hamper the effectiveness of the coated pancreatic enzymes, as already mentioned. Acid lipases of non-pancreatic origin, such as gastric lipases or lipases from other species, have been shown to be present in animal models with CF and could be used therapeutically. So far no trial using acid lipases in CF patients has been published.

UROGUANYLIN

Uroguanylin is an intestinal peptide hormone that may regulate epithelial ion transport by activating a receptor, guanylyl cyclase, on the luminal surface of the intestine. In mice this drug stimulated ion transport mainly in the proximal duodenum and in the cecum and less in the jejunum, ileum, and distal colon. Acidification of the lumen of the proximal duodenum (pH 5.0–5.5) enhanced the stimulatory action of uroguanylin. Experiments, using pH-stat titration, revealed that uroguanylin stimulates serosal-to-luminal HCO_3^- secretion. In CFTR knockout mice, the duodenal response to uroguanylin was markedly reduced, but not eliminated, despite a similar density of functional receptors. It was concluded that uroguanylin is most effective in acidic regions of the small intestine, where it stimulates both HCO_3^- and Cl^- secretion primarily via a CFTR-dependent mechanism. In vivo studies in mice have demonstrated that orally administered uroguanylin resists luminal proteolysis and stimulates net intestinal fluid accumulation. On the basis of its action to increase HCO_3^- secretion, oral uroguanylin may be potentially useful as a pharmacological agent for the prevention and treatment of duodenal ulcers and probably to improve function of pancreatic enzymes in CF. [69]. However no trial with this peptide hormone has been published in CF.

ALTERNATIVE INTESTINAL BICARBONATE PRODUCTION

It has been demonstrated that human duodenal enterocytes contain at least three acid/base transporters: a Na^+/H^+ exchanger that serves to extrude acid, a Na-bicarbonate cotransporter that functions as base loader, and a $\text{Cl}^-/\text{HCO}_3^-$ exchanger that operates as a base extruder [70]. These alternative pathways can be activated by carbachol and by heat-stable toxin of *Escherichia coli*. [71]. Until now no attempts concerning the use of such therapies in CF patients have been reported.

ANTACIDS

One of the most commonly used self-prescribed drugs are antacids, consisting of calcium carbonate and magnesium – aluminium salts in various compounds or combinations. The effect of antacids on the stomach is due to partial neutralization of gastric hydrochloric acid (HCl) and inhibition of the proteolytic enzyme, pepsin. Antacids have been used in CF, but no beneficial effect was observed. Therefore antacids alone are insufficient in improving fat malabsorption in CF [72].

H₂ RECEPTOR ANTAGONISTS

The need of antacids for intestinal ulcer related diseases was reduced after the introduction of H₂ receptor antagonists. These drugs were investigated in CF and have shown to be only slightly effective in improving malabsorption when compared to PPI [72, 73]. In the tables 1.1. and 1.2. we have included trials with this type of medication.

PROTON PUMP INHIBITORS (PPI)

The most effective agents for suppressing gastric acidity are PPIs, which are the treatment of choice for many acid-related GI tract conditions [74]. These drugs potentially inhibit acid secretion from gastric parietal cells by irreversibly inhibiting activity of the H^+/K^+ ATPase (proton pump). They are predominantly inactivated by the 2C19 isoform of the hepatic cytochrome P450 mixed function oxidase system. Some individuals present several different mutations for this system and will consequently metabolize PPIs more slowly resulting in higher drug levels. Therefore, the efficacy of acid suppression by standard doses of a PPI varies in the general population [75]. This might also explain the differences in potency between several PPIs, where lansoprazole and rabeprazole provide earlier and better symptom relief than omeprazole and pantoprazole. [76]. Rabeprazole is less dependent on 2C19 for metabolism and therefore the most potent [77]. Small molecular changes can result in important improvements as has been shown for esomeprazole [78]. PPIs do increase the pH of the proximal duodenum for a prolonged period, especially shortly after meals, and based on superior efficacy profiles they are the drugs of choice in patients with peptic ulcer disease, gastroesophageal reflux disease (GERD) and Zollinger-Ellison syndrome [79]. Also in CF patients they have been used with success [93,94,98].

TABLE 1.1. Effect of short-term (<3 months) use of gastric acid inhibitors in CF

1 st Author	Number cases	Age (yr)	Type of study	Pancreatic enzyme	Intervention	Duration	Effect on steatorrhea	Comments
Cox '79 [81]	10	6-27	Prospective Crossover Open	Cotazym Viokase	Cimetidine 150-200 mg/day	1 week	Fat excretion ↓ N excretion ↓	No diet evaluation
Boyle '80 [82]	8	12-25	Prospective Crossover	Viokase	Cimetidine 300 mg/day	5 days	Fat absorption coefficient ↑ Fat excretion ↓ Fecal weight ↓	No diet evaluation
Durie '80 [83]	15	10-17	Randomized Prospective Crossover	Cotazym	Cimetidine 20 mg/kg/day	7 days	Fat absorption coefficient ↑ Fat excretion ↓ N excretion ↓ Fecal weight ↓	No effect if fat intake >120g/day
Hubbard '81 [84]	8		Randomized		Cimetidine 900 mg/day	6 weeks	Fat excretion ↓ FFA en lipids ↑ Gastric acid ↓	
De Bievil '81 [85]	13			Pancrease	Cimetidine 2 doses		Fat and N excretion: NS	
Gow '81 [86]	10	6-13	Prospective Crossover	Pancrease	Cimetidine 20 mg/kg/day	14 days	Fat and N excretion, Fecal weight: all NS	
Schöni '81 [87]	10	11-17	Randomized Prospective Open	Eurobiol	Cimetidine 600 mg/m ² /day	6 days	Fat and N absorption coefficient: NS	

N excretion = fecal losses of Nitrogen bound products; FFA = free fatty acids; NS = not significant

TABLE 1.1. CONTINUED. Effect of short-term (<3 months) use of gastric acid inhibitors in CF

1 st Author	Number cases	Age (yr)	Type of study	Pancreatic enzyme	Intervention	Duration	Effect on steatorrhea	Comments
Stead '88 [88]	14	adults	Open Randomized Crossover	Pancrease Non/EC pancreatin	Non-EC with Cimetidine vs EC pancreatin alone	28 days	Fat excretion and Fecal weight: NS	Compared non- / EC pancreatin
Cleghorn '88 [89]	11	2-17	Prospective Open	Pancrease	Misoprostol 400 mg/day	1 week	Fat excretion ↓ Fat absorption coefficient NS	Fat intake not controlled
Robinson '88 [90]	22			Pancrease	Misoprostol 400 mg/day		Fat excretion ↓ Fat absorption coefficient ↑	Only in patients fat excret. >10%
Robinson '90 [91]	17		Doubleblind Placebocontr. Crossover	Pancrease	Misoprostol 400 mg/day	6 weeks	Fat excretion ↓ Fat absorption coefficient ↑	Only in patients fat excret. >10%
Heijerman '90 [92]	8		Doubleblind Placebocontr. Crossover	Pancrease	Ranitidine 2x150 mg Enprostil 2x 35 mg	14 days	Ran. > Enpr. Fat excretion ↓ Fecal weight ↓	
Heijerman '91 [93]	9		Doubleblind Crossover	Pancrease + doubling	Omeprazole 1 dd 20 mg	14 days	Fat excretion ↓	Only in Pancr.2x + Omeprazole
Heijerman '93 [94]	11	20-42	Doubleblind Placebocontr Crossover	Pancrease	Omeprazole 1 dd 20 mg	14 days	Fat excretion ↓	

EC = enteric-coated; NS = not significant; Pancr.2x = double Pancrease dose

TABLE 1.2. Effect of long-term (≥3 months) use of gastric acid inhibitors in CF

1 st Author	Number cases	Age (yr)	Type of study	Pancreatic enzyme	Intervention	Duration	Effect on steatorrhea	Effect on nutritional status
Schöni '84 [95]	38	Mean 13	Prospective Doubleblind Randomized	Pancrease	Cimetidine 600 mg/m ² /day	4 months	Plasma lipids and lipoprotein: NS	W/H: NS Skinfolds NS TLC/TGV: NS Raw/SGAW ↑
Chalmers '85 [96]	17	5-19	Placebocontr Randomized Crossover	Pancrease	Cimetidine	6 months	Fat excretion ↓ Fecal weight and N excr.: NS	W/H: NS Skinfolds NS Clinical score NS
Carroccio '92 [97]	10	7-18	Doubleblind Crossover	Pancrease	Famotidine 1 mg/kg/day	6 months	Fat absorption Coefficient ↑ Fecal weight ↓	W/H ↑ Clinical score: NS
Tran '98 [98]	15	3-22	Prospective Crossover	Pancrease	Lansoprazole 15 mg/kg/day	3 months	Fat absorption Coefficient ↑	W/FM/BMC ↑ Skinfolds ↑ FFM: NS

NS = not significant; W/H = weight and height; TLC/TGV = total lung capacity and thoracic gas volume; Raw/SGAW = ademweerstand/specifieke conductantie van de luchtwegen; N. Excr. = fecal losses of Nitrogen; BMC = bone mineral contents; FFM = fat-free mass

VAGAL NERVE RESECTION

Although vagal nerve resection could have a beneficial effect on luminal pH, by decreasing gastric acid production, the vagal nerve also stimulates hepatic, pancreatic and duodenal mucosa bicarbonate production. The latter was demonstrated by electrical vagal stimulations in pigs [80]. As vagal nerve resection would decrease the necessary non-pancreatic bicarbonate secretion. Due to its invasiveness we consider this technique not as a real option in CF patients.

SUMMARY

In CF several studies concerning the use of duodenal pH increasing therapy, have been performed. Table 1.1. shows the short term studies (<3 months) of gastric acid inhibitors, mainly H₂ blockers, on malabsorption. These studies were too short to evaluate the effects on the nutritional status. Table 1.2. shows the few long-term studies (≥ 3 months) of gastric acid inhibition, assessing nutritional status. More studies assessing nutritional interventions and/or trials with different pancreatic enzymes have been performed. The possible disadvantages of the use of PPIs, especially in children with CF, are discussed in the next section.

1.7. PROS AND CONS FOR THE USE OF PPI IN CHILDREN WITH CF

POTENTIAL BENEFICIAL ROLE OF PPI IN CF

In the following sub-sections several considerations for the use of PPI in CF are described.

Fat malabsorption: The major effect of PPIs is a decreased need of pancreatic bicarbonate for buffering gastric acid entering the duodenum. This should result in an earlier opening of the enteric-coated capsules, less denaturation of enzymes, better lipolysis, and an increase in the concentration of ionized bile salts. All these factors will improve fat absorption.

Absorption of fat soluble vitamins: Besides the fecal loss of lipids, steatorrhea will also result in increased losses of several hydrophobic vitamins. At risk for deficiencies are the fat-soluble vitamins A, D, E and K. Despite supplementation with standard multivitamins and pancreatic enzymes, the occurrence of persisting fat-soluble vitamin deficiency is relatively common in CF [99]. Vitamin K deficiency is a typical example. This vitamin is an important co-factor in the carboxylation of osteocalcin (OC), an important protein in bone formation. In CF patients a greater fraction of OC seems to be under-carboxylated. Recent work has highlighted the association of reduced bone density with the prevalence of vitamin K deficiency. Deficiency of both vitamins D and K could explain the decrease in bone density in CF patients before their puberty, reaching a lower peak bone mass as adolescent [100, 101]. By decreasing fat excretion, the addition of a PPI might improve the vitamin status of CF patients.

Gastroesophageal reflux disease (GERD): Another important issue is GERD, which has been shown to occur frequently in CF. In untreated infants with CF, one in five had pathological GERD and one in four had increased acid reflux time, while only 54% showed normal oesophageal pH recordings. Pathologically increased reflux was found to be present before radiological lung disease was established. Apart from frequent vomiting, no useful clinical symptoms of pathological reflux were found [102]. By inhibiting gastric acid production, acid reflux will decrease. This acid reflux can induce inflammation in the ENT area [103]. It might also contribute to pulmonary inflammation and therefore almost half of the young CF patients could benefit from the use of PPIs.

Increased intestinal permeability (IP): As already mentioned IP is severely increased in CF, and abnormalities related to the presence of pancreatic dysfunction in CFTR may contribute to this increase of IP. Especially the associated decrease in ductular bicarbonate secretion, resulting in a decreased duodenal buffering capacity and therefore in an acid luminal pH, could be responsible for this [104]. Acid intestinal pH could either directly damage tight junctions between enterocytes or be responsible for late opening of the enzyme coating in the small bowel, resulting in elevated luminal proteolytic activity, which will also damage enterocytes and increase IP. Therefore the use of PPIs, by increasing the intestinal pH, may have a beneficial effect on intestinal permeability.

ADVERSE EVENTS BECAUSE OF THE USE OF PPIs

Potential areas of concern regarding long-term PPI use have included:

Carcinoid formation and development of gastric adenocarcinoma: Prolonged PPI use may lead to enterochromaffin-like cell hyperplasia, but it has not been shown to increase the risk of carcinoid formation. There is no documentation of long-term PPI treatment hastening the development or the progression of atrophic gastritis to intestinal metaplasia and gastric cancer, although long-term studies are required to allow definitive conclusions. It seems that patients with *Helicobacter pylori* infection could be more prone to developing carcinoid formation [105].

Bacterial overgrowth and enteric infections: Gastric bacterial overgrowth is related to the degree of acid suppression, but clinically important sequelae, such as a higher rate of gastric adenocarcinoma, have not been seen. The risk of enteric infection may increase with acid suppression, although this does not seem to be a common clinical problem with prolonged PPI use [105].

Malabsorption of minerals, and vitamins: The absorption of vitamin B₁₂ may be decreased when gastric acid is markedly suppressed for prolonged periods (e.g. Zollinger-Ellison syndrome) due to an impaired release of vitamin B₁₂ from food in a non-acid environment. The malabsorption of crystalline vitamin B₁₂ observed in CF is due to a mechanism independent of either intrinsic factor secretion or intrinsic factor biological activity. The absorption of crystalline vitamin B₁₂ in CF is corrected in most

patients by administration of pancreatic supplements [106]. Anyhow PPIs do not promote the development of pernicious anemia [107].

THE LONG-TERM USE OF PPIs IN CHILDREN

Knowledge about this class of drugs in pediatrics is still fragmentary, but the reported pharmacokinetic and clinical data indicate that they are suitable for use in children. While the short-term risk of complications appears to be minimal, the long-term tolerance requires careful monitoring because of the potential consequences of prolonged inhibition of acid secretion [108]. Current evidence suggests that prolonged gastric acid suppression with proton pump inhibitors rarely, if ever, produce adverse events. Nevertheless, continued follow-up of patients taking proton pump inhibitors for extended periods will provide greater experience regarding the potential gastrointestinal adverse effects of long-term acid suppression.

CONCLUSION ABOUT PROS AND CONS OF PPIs IN CHILDREN

In the present state of knowledge the potential advantages of using PPIs in pancreatic insufficient CF patients seem to outweigh the potential drawbacks. Awareness for unknown adverse effects after long-term use of these type of drugs is particularly required in children.

1.8. AIMS OF THE STUDY

In view of the persisting malabsorption in pancreatic insufficient CF patients despite enzyme supplementation and while awaiting the potential breakthrough of gene therapy, we aimed at improving the nutritional condition of our patients, by using PPIs in order to increase luminal pH and consequently decrease their persisting malabsorption. We assessed not only the effects on fat malabsorption of such a treatment but tried also to elucidate the following questions:

- ❖ Is there a relationship between body composition and lung function?
- ❖ Could PPIs modify the gut permeability?
- ❖ Could PPIs modify airway hyperresponsiveness?
- ❖ Could PPIs modify the essential fatty acid deficiency?
- ❖ Is vitamin B₁₂ deficient after prolonged use of PPIs?

The successive chapters in this thesis will give the study design, aims, results and interpretations of these questions. As the complete set of studies is of an observational design, definite conclusions can not been made. Therefore at the end of this thesis we will sum up the results, delineate a role for the use of proton pump inhibitors in young CF patients, but we can not advise the routine use of PPIs in children with CF.

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CHAPTER TWO

Trial Design: An open study to assess the effects of one year lansoprazole treatment on nutritional status, body composition, lung function and intestinal permeability in children with Cystic Fibrosis

Han Hendriks

AN OPEN STUDY TO ASSESS THE EFFECTS OF ONE YEAR LANSOPRAZOLE TREATMENT ON NUTRITIONAL STATUS, BODY COMPOSITION, LUNG FUNCTION AND INTESTINAL PERMEABILITY IN CHILDREN WITH CYSTIC FIBROSIS

An observational study to assess the effects of a pharmacologic modulation of gastric acid secretion on biological variables in children with Cystic Fibrosis. This chapter describes the aims, the subjects and the methods used in our studies performed in young CF patients which are presented in the chapters 4 through 8.

2.1. BACKGROUND AND AIMS OF THE COMBINED STUDY

In a previous, short term study we evaluated the effect of a proton-pump-inhibitor (PPI), lansoprazole (Prezal[®]), on fecal fat excretion and body composition, in children with Cystic Fibrosis (CF). One of the main issues in CF disease is exocrine pancreatic insufficiency, resulting in a shortage of both enzymes and bicarbonate secretion. The lack of pancreatic bicarbonate results in a poor duodenal neutralization of gastric acid and a duodenal pH below the pH optimum (5.5) of pancreatic enzymes activity. Fat digestion will be hampered, resulting in steatorrhea and malnutrition. Based on the results of our “pilot” study, increasing duodenal luminal pH through the use of a PPI could be expected to decrease fecal fat excretion and enhance body composition. Additional effects that might be expected are a decrease of the abnormally elevated intestinal permeability (IP) characteristic of CF patients, and secondarily also improvements of the essential fatty acids status and lung function. Finally PPIs might induce side effects, like vitamin B₁₂ deficiency. Therefore, the aim of this study was to investigate whether or not during a pharmacological intervention with PPIs in children with CF any of these predictable effects occur. Secondly, this study was also intended to provide data to perform a power analysis for possible future, placebo-controlled studies. For both purposes an observational study, in children with CF and ongoing pancreatic insufficiency, was performed.

2.2. SUBJECTS

STUDY POPULATION

Children with a confirmed diagnosis of CF involving the respiratory tract and gastro-intestinal tract were eligible for the study. All children received standard therapy, including several pharmacological interventions (described in 2.3) and hyperalimentation (described in 2.4), as part of their regular management before the start of the study. A total of 14 patients entered the study, all completing a one year treatment protocol. All but one patient continued the use of the PPI for 3 years allowing us to study long term effects. Approval of the medical ethical committee of the university hospital of Maastricht and informed consent of the parents and patients older than 11 years was obtained.

TABLE 2.1. Nutritional daily recommendations for children with CF

Energy	RDA $\times$ (1.2-1.5)
Protein	RDA $\times$ (1.2-1.5)
Fat	35-40 % of energy
Essential fatty acids	5 % of energy
Sodium chloride	Infants 1 g/d, children 3-6 g/d (compensation for sweat losses)
Fat soluble A	2500-5000 IU in water-soluble form
Vitamins: D	800 IU
E	5-10 mg/kg (max. 300 mg) tocopheryl acetate
K	not defined (probably 1 mg/kg/day)
Water soluble B group	RDA $\times$ 2
Vitamins: B12	RDA unless ileal resection
C	RDA $\times$ 2
Iron, Zinc	RDA (suppletion only if deficient)

RDA= recommended daily allowance (for healthy children)

CRITERIA WHICH ALL HAD TO BE FULFILLED, BEFORE INCLUSION.

1. The diagnosis of CF had to be confirmed by sweat test according to the method by Gibson and Cooke [1] (chloride of more than 60 mMol/L) or by DNA analysis.
2. The patients had to be able to perform lung function measurements adequately and their pre-study FEV₁, in the preceding year, had to be over 70% of mean values for normal children with the same height and sex (predicted).
3. The patients had to demonstrate ongoing pancreatic insufficiency, despite an intake of at least 2000 IU lipase/kg body weight, as indicated by a pre-study stool acid steatocrit (an index of fecal fat excretion) over 25% in the preceding year.
4. The patients had to receive the recommended daily nutritional requirements for CF patients, as outlined in table 2.1.
5. The patients had to receive the recommended medical and physiotherapeutical treatment and they should not have used PPIs in the preceding year.
6. No participation in any trial was allowed within the previous 3 months.

CRITERIA LEADING TO EXCLUSION FROM THE STUDY

1. Hospital admission in the 3 months preceding the study or administration of (oral) antibiotics at home during one month preceding the study (infectious exacerbation).
2. Other metabolic disorders including diabetes mellitus and thyroid disease, which might confound interpretation of the study results or clinically significant cardiovascular, renal, central nervous or musculo-skeletal system disorders.
3. Severe malnutrition requiring daily enteral (by nasal tube or gastrostomy) or parenteral nutritional supplementation, in addition to that described in table 2.1.
4. Considered by the investigator to be unsuitable for the study for reasons of non-compliance or inability to follow instructions correctly.

TABLE 2.2. Patient characteristics at start (N=14)

Item		Mean	SD	Range
Age	(year)	9.74	3.38	4.6 / 17.2
Height	(cm)	133.9	16.6	107.0 / 162.3
	(SDS)	-1.068	0.590	-2.166 / +0.003
Weight	(kg)	27.7	9.9	15.5 / 52.3
	(SDS)	-1.270	0.739	-2.393 / +0.272
Acid Steatocrit	(%)	33.3	17.7	5.4 / 65.7
FEV ₁	(% predicted)	83.9	17.2	58 / 112
FVC	(% predicted)	86.6	13.0	64 / 109

2.3. MEDICAL TREATMENT

During one year lansoprazole (Prezal[®]) was administered at a dose of 15 mg once daily, 30 minutes before breakfast. All other medication used before the start of the study was continued in the same dose and frequency. All patients used nebulized saline daily, with or without mucolytics, while four also used nebulized DNAse (Pulmozyme[®]). All patients performed regularly sputum mobilization exercises as instructed and controlled by their physiotherapist. Three patients were on ursodeoxycholic acid, while no patient was on daily antibiotics. New medication was permitted, if necessary, according to standard protocols for CF care. Infectious exacerbations, as occurred in 4 children once, were treated according to the usual protocol, with antibiotics for at least 2 weeks. During the whole study year, only one child was admitted because of complaints of his previously demonstrated gastro-intestinal fistula.

2.4. DIET AND PANCREATIC ENZYMES

A free but constant diet, similar to that used before the study, according to recommendations (table 2.1.), was prescribed to each patient during the study. The only difference was that during the study the dietary intake was analyzed extensively, every 6 months, instead of once a year. For that purpose at start and after 6 and 12 months, parents and their children held, for three consecutive days, a detailed record of both nutrient intake and pancreatic enzymes doses. This was analyzed and computed by one dietary consultant. For each 6 months period and for each different meal, changes in nutrients, energy and lipase intake were analyzed.

2.5. STUDY SCHEME

(Months)	0	3	6	9	12	24	36
<div></div>							
Visits							
Clinical evaluation							
+ Anthropometry	x	x	x	x	x	x	x
Dietary evaluation	x		x		x	x	x
Body composition							
DEXA	x		x		x		x
Resting energy expenditure							
Ventilated Hood	x	x	x	x	x		
Gastro-intestinal function							
3 sugars permeability test	x				x		
Fecal fat content	x	x	x	x	x	x	x
Urinary NOx/creatinine-ratio	x	x	x	x	x		
Lung function							
- F/V-curves, bodybox	x	x	x	x	x	x	x
- PEF	4 times daily - 1 week/month						
Laboratory assesments							
Fatty Acids status							x
Vitamin B ₁₂ -homocysteine							x

2.6. METHODS

All patients were already followed by regular 3 monthly out-patient clinic visits. Additional investigations required for the present study were extra lung function tests, urine sampling for NO oxidation products, DEXA scans, two weekly fecal samples and the 3-sugars test. Also the PEF measurements performed at home during one week every month and one extra blood sampling for essential fatty acids status, vitamin B₁₂ and homocysteine blood levels were on top of routine assesments. Altogether at the start and at one year the out-patient clinic visits and required assessments lasted 6 hours, at 6 months they took 4 hours and at 3 and 9 months only 2 hours. The assessments at 2 and 3 years were obtained during routine out-patient clinic visits, extended once yearly with check-up according to our regular protocol. As all investigations used in the present study were non-invasive, it was not unexpected that there were no missing data in the first year. We only lack the data of our oldest patient at 36 months.

ASSESSMENT OF ANTHROPOMETRIC PARAMETERS

Clinical evaluation: Every 3 months the patients were seen at the out-patient clinic for a routine systematic physical examination covering all body systems, including ENT, respiratory, gastro-intestinal, cardiovascular, neurological, muscles, skeletal, and skin. All medication was checked and prescribed for the following 3-month period.

Weight and Height: At visits in the out-patient clinic, weight and height, as mean of three separate measurements, was assessed, by one single investigator (Hendriks), with the subjects barefoot and dressed in underwear. The same instruments, a Harpenden stadiometer (Holtain Ltd., Crymych, D Y Fed. UK) and an electronic weight scale (Seca Delta, Model 707) were used during the study period. Body mass index (BMI) was computed as weight/height² in (kg/m²). Weight, height and BMI are expressed as standard deviation scores (SDS) for age and sex. We used Dutch reference values, as recorded by Gerver and de Bruin [2,3]. Expression in SDS changes is a prerequisite for pediatric studies, as absolute changes in weight and height with time are age and sex dependent and would be uninterpretable in a group of children of different ages and sexes.

BODY COMPOSITION BY TOTAL BODY DUAL ENERGY X-RAY ABSORPTIOMETRY (DEXA)

This is a non-invasive method, which is used for an easy and rapid (<30 minutes) evaluation of the body composition and bone mineral status. During one assessment total body fat (TBF), fat mass (FM), fat free mass (FFM), bone mineral content (BMC) and bone density (BMD) can be measured. A pediatric software program (Paediatric Beta Software: Lunar 1.5) was used. The radiation dose of this examination is very low (<5 mRem) when compared with the standard yearly background radiation exposure (82 mRem/year). This method is considered to be the method of choice for measuring body composition in young children. Because the children are in a normal growth process, results were compared with those of a large (n=403), normal Dutch reference group, matched for age and sex, and expressed as SDS [4].

RESTING ENERGY EXPENDITURE (REE)

Basal metabolic rate or REE was measured by indirect calorimetry (ventilated hood) and compared to Schofield's age-related normal values for children [5,6]. The principle of a ventilated hood system is that a stream of air is forced to pass across the face of a subject and mixes with the expired air, which is collected by a transparent plexiglas hood (30 L) placed over the subject's head. The rate of REE was calculated by determining the amount of air flowing through the hood and by measuring the oxygen and the carbon dioxide concentrations in the in and out coming air. The assessments were performed after an overnight fast in the morning, during tidal breathing in supine position for 15 minutes, at complete rest.

GASTRO INTESTINAL FUNCTIONS

Fecal Fat Content: Steatorrhea was monitored by measuring the fecal acid steatocrit as previously described [7]. According to this method results are measured as fat / fat + solid layer and expressed in percentage. Fecal acid steatocrit has been shown to correlate well with chemical analyses of fecal fat ($r = 0.83$). We chose this method in order to monitor intensively and easily fecal fat content, allowing us to obtain regular recordings during the study. Before the start (3 samples, within 2 weeks) and every two weeks during the entire study a small stool sample was taken, placed in a container, wrapped in aluminum foil, stored in the refrigerator at home at -18°C and taken to the hospital during visits. The upper normal limit for acid steatocrit in children older than 0.5 years is 20%, as described elsewhere [8,9].

Intestinal Permeability (IP): IP was measured before and after one year treatment with lansoprazole, by the 3-sugars permeability test (3-SPT), according to Miki et al [10]. After an overnight fast, the first urine of the day was discarded and the patients ingested 100 ml of an isotonic test solution (osmolality 330 mOsm/L) containing 1 gram Mannitol (M), 1 gram L-Rhamnose (Rh) and 5 gram Lactulose (L). Only water intake was allowed after the first half-hour of this study, while food and sugar containing fluid intake were withheld for 4 hours. Urine was collected for 5 hours for determination of these 3 sugars by high-performance liquid chromatography. Results were expressed as percentage of intake. Sugar excretion ratios were determined by dividing the percentage of urinary recovery of the large sugar (L) by that of the smaller sugars (M + Rh). This simple and non-invasive test is a reproducible index of IP in children [11]. The use of ratios eliminates potentially confounding factors such as delayed gastric emptying, delayed or rapid gut transit, or impaired renal clearance.

LUNG FUNCTION

An experienced lung function technician performed the lung function tests, using a Jaeger Masterlab pneumotachograph and bodybox. Calibration of the apparatus was performed twice daily. Only lung function data with FVC variability of less than 5%, on 3 consecutive maneuvers, were accepted. Lung function parameters were expressed as % of predicted (100% is the mean of normal children, with the same height and sex) according to the Zapletal reference values [12]. The following lung function parameters were used:

Static and Dynamic lung volumes

Flow Volume curves: (particularly)

- ◆ Forced Expiratory Volume in 1 second (FEV_1)
- ◆ Forced Vital Capacity (FVC)
- ◆ Peak Expiratory Flow (PEF)

Single body box:

- ◆ Total Lung Capacity (TLC)
- ◆ Residual Volume (RV)
- ◆ Intra Thoracic Gas Volume (ITGV)
- ◆ (specific) Airway Resistance and Conductance (R_{aw} , sR_{aw} , G_{aw} , sG_{aw})

Respiratory muscle function: As representative of respiratory muscle function, maximal mouth pressures were measured at residual volume (RV) for maximal inspiratory pressure (PI_{max}) and at total lung capacity (TLC) for maximal expiratory pressure (PE_{max}), according to the method of Black and Hyatt [13]. As there are no normal values for children, the exact data are given.

Peakflow (PEF) variability and Bronchial Lability (BL): In asthmatics, bronchial hyper-responsiveness (BHR) can be increased due to both recurrent infections or gastro esophageal reflux (GER). Because of a different patho-physiology in CF and because of drawbacks of the PEF measurements, it is perhaps better to describe it as bronchial lability (BL). Although GER is known to be present in many (young) CF patients, the existence of BL and the effect of PPI on this BL has never been examined in these patients. Daily PEF variability is an easy and commonly used lung function assessment, especially in normal daily life conditions. In the present study PEF was measured at home four times daily, using a "Personal Best" PEF meter (Respironics HealthScan Products, Europe, Belgium). Instruction was given about the correct technique and reading of the PEF meter. In order to detect an incorrect use, three consecutive maneuvers were performed and recorded, while only the best of these 3 measurements was used for data management. In order to measure changes in BL, several PEF characteristics were calculated from the diary records performed for an one week period every month, during the first year of the study [14]:

- ◆ Mean diurnal variability (in L/min), 7 days with 4 data per day.
- ◆ Maximal diurnal variability: the difference between the worst (morning) and the best (evening) PEF of that week (expressed as % of recent (that week) best PEF).
- ◆ Minimal morning PEF (expressed as % of recent best PEF).
- ◆ The difference between the best morning and the best evening PEF (in L/min).

MUCOSAL INFLAMMATION

Mucosal inflammation has been shown to be present in CF patients [15,16]. Urinary nitric-oxide products (NO_x) are markers of cytokine-induced increase of mucosal NO-synthesis. Like others we could show that NO_x are increased in acute celiac disease [17,18]. We asked the patients to collect their urine before and at 3 months intervals during this study, in order to evaluate this marker in CF. Twenty-four hours prior to a visit, the CF patients collected urine, after using a nitrate-poor diet for 48 hours, before and during the collection period. Patients were free of viral illness or CF exacerbation during the collection period. Nitrite and nitrate content were determined using a Nitrate/Nitrite colorimetric assay kit (Cayman Chemical Company, Ann Arbor, MI, US). This test uses the Griess reagent assay, which is more accurate than

the HPLC method [19]. Urinary creatinine was also determined in order to compute the NOx / Creatinine ratio. The data collected in our CF patients were compared with results of urine samples of 25 normal children (mean age 9.0 (± 4.8) years).

2.7. SPECIAL LABORATORY MEASUREMENTS

Blood was not only taken for routine blood analyses, but also for special assessments. The following investigations were performed.

Fatty Acids: The fatty acid (FA) status was evaluated in 20 CF patients (11 with PPI and 9 without PPI) and in 10 children, submitted to minor surgical procedures (controls). The fatty acid profiles were determined in plasma and red blood cell phospholipids as previously described [20]. Briefly, lipid extracts were prepared from both plasma and RBC samples, and the phospholipid fraction was separated. Then the phospholipid FA were hydrolyzed and transmethylated. The composition of the obtained FA methyl esters was determined by capillary gas chromatography using a polar column, with helium as carrier gas and at a pressure of 130 kPa. The amount of each fatty acid was quantified by the amount recovered of the internal standard (dinonadecanoyl [19:0]-phosphatidylcholine) added to each sample.

Vitamin B₁₂ and Homocysteine: Vitamin B₁₂ and homocysteine blood levels were measured in 20 CF patients (11 with PPI and 9 without PPI) and in 10 children, submitted to minor surgical procedures (controls). Total homocysteine concentration in plasma was measured by HPLC, as previously described [21], and vitamin B₁₂ by a competitive protein binding assay (Perkin-Elmer, Turku, Finland).

2.8. STATISTICS

As our aim was to evaluate possible effects of PPIs, all measured parameters were analysed in two ways. Firstly, in order to maximize the chance to find a possible effect, we compared the mean of the whole intervention period to the pre-trial (baseline) values whenever possible. In that way, the power of the test is optimized, since measurement errors and within-subject variation tends to cancel out. A second advantage of this method is that we avoided doing 4 different tests for essentially the same hypothesis, which would inflate the type-I error probability beyond the nominal 5%. By performing this procedure a Bonferroni correction is not necessary and, according to Altman, this is advised as one of the five possibilities to avoid a repeated measurements design [22]. Secondly, as this procedure will show a less detailed picture, we also looked for trends within the study period. Furthermore, by using the non-parametric Wilcoxon signed ranks test, we avoided to make the assumption of normality, which is untestable with only 14 subjects. Differences with respect to baseline were considered significant when the two-sided *P*-value was 0.05 or less. The

statistical package SPSS (version 9.0 and 10.0 for Windows, 1998) was used for data management and analysis.

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CHAPTER THREE

Longitudinal growth of children with Cystic Fibrosis: a longitudinal retrospective study

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3.1. ABSTRACT

Background: Studies of growth in children with Cystic Fibrosis (CF) used cross-sectional data and showed a retarded growth pattern in CF. We plotted the growth data of 27 children with cystic fibrosis longitudinal, in order to define growth patterns in children with CF.

Methods: Mean Standard Deviation Scores (SDS) and 95% confidence intervals for height, weight and weight-for-height were plotted for age intervals. Patients were followed up to the age of 120 months. Non-parametric two-samples tests (Mann-Whitney-U-test) were used in order to compare SDS results between boys and girls at various time intervals.

The Pearson correlation coefficient was used for studying the relationship between catch-up growth and the severity of the initial growth failure.

Results: Compared to normal children, both boys and girls initial showed a severe decrease in mean SDS for weight (boys -3.28; girls -2.84), height (boys -2.88; girls -1.84) and weight-for-height (boys -1.11; girls -1.02). Catch-up growth in CF patients occurred in the first year after diagnosis and resulted in normalising of the SDS for each parameter at the age of two years. A strong correlation was found between catch-up growth for height, weight and weight-for-height and the severity of the initial growth failure. A slight decrease in both mean weight and height SDS occurred when children grew older while weight-for-height remained normal.

Conclusions: Growth patterns show stunting more than wasting, making the use of weight-for-age growth charts preferable to that of weight-for-height in these patients. These data further support the need for CF screening programs as initial growth was in 80% severely decreased (less than - 2 SD) and early intervention could prevent stunting in these patients.

3.2. INTRODUCTION

Cystic Fibrosis (CF) as a chronic disease is a known cause of growth retardation [1-13]. In CF, growth failure may result from insufficient energy intake, malabsorption and increased catabolism [4,5,9-13]. After making the diagnosis and starting therapy, catch-up growth is known to occur and has been achieved by treating the two main underlying disease entities malnutrition and chronic airway infection. Therefore, besides antibiotics, nutritional support with high-energy diets and by optimising pancreatic supplementation is very important [1-7,10,11,13]. Early detection by screening and treatment in CF centres further improved the outcome of CF patients. In these settings, CF patients experienced significantly fewer and shorter hospital stays in the first year of life and malnutrition in many patients was prevented [7,11]. Catch-up growth for height in children with CF has been reported to be incomplete [3-5,7,9,12,13] and may be also one of the reasons why CF patients start in about one year later with their pubertal growth spurt [14]. Catch-up growth for weight has also been reported to be incomplete, except for one study where weight and height catch-up growth was reported to be complete within 12 months [6].

When studying growth disturbances in childhood the indices of stunting and wasting are used. Stunting and wasting were previously defined as chronic and acute malnutrition respectively. The present definition of wasting is that of a deficit in weight-for-height, while stunting is defined as being a deficit in height-for-age [15]. The first aim of the present retrospective study was to evaluate long term longitudinal growth of children with CF. The second aim was to study the catch-up growth pattern in these patients. Answers to both of these questions could help resolve the dilemma "to screen or not to screen" for this condition.

3.3. SUBJECTS AND METHODS

PATIENTS

Follow-up growth data of 27 CF patients (11 females) were available for our study. The patients were born between 1977 and 1996, therefore patients had variable follow-up periods. Only patients diagnosed before the age of 4 months were included and diagnosis was confirmed by repeated sweat tests. Mean present age (SD) was 11.7 (4.0) years for girls and 9.3 (5.2) years for boys. Twenty-four patients were pancreatic insufficient, while three were pancreatic sufficient.

METHODS

Weight and height measurements were obtained from the charts of the out-patient department of Pediatrics of the University Hospital Maastricht. The children were measured using standard anthropometric techniques, using the same instruments, a Harpenden stadiometer (Holtain limited, Crymch, D Y Fed., UK) and an electronic weight scale (Seca delta, modell 707). The data were expressed as SDS, making use of the normal pediatric Dutch reference values, collected by Gerver and de

Bruin [16]. SDS is defined as the number of standard deviations from the mean. These SDS were calculated for height, weight and weight-for-height. The patient mean and 95% confidence interval SDS were plotted for different age periods. The study period was divided in six months periods up to 120 months, except for the first year of life where three periods of each 4 months were used.

Growth data have been studied for girls and boys separately and, due to the lack of significant sex differences, as well as the use of SDS, have been pooled together in the figures.

Catch-up growth was defined as the difference between the best SDS, between 18 and 36 months of age, when SDS had most often normalised, and the worst SDS at diagnosis, at the age of 0 to 8 months. The relationship between catch-up growth and the severity of growth failure at the start was evaluated by correlating catch-up growth with the worst initial SDS.

STATISTICAL ANALYSIS

Parametric tests were used for the calculation of mean values and confidence intervals. A significant difference was considered to exist between the reference population and our data when the mean value of the data set differed from 0 at the 0.05 confidence level. Non-parametric two-sample tests (Mann-Whitney-U-test) were used in order to compare SDS results between boys and girls at various time intervals. The Pearson correlation coefficient was used for studying the relationship between catch-up growth and the severity of the initial growth failure. The SPSS 7.5 for Windows statistical package was used for data management.

3.4. RESULTS

Mean SDS for height, weight and weight-for-height for the different age periods are shown in table 3.1. The initial mean height SDS were evidently decreased for

TABLE 3.1. Height, weight and weight-for-height SDS for female and male CF patients.

Age periods	Sex	Height SDS		Weight SDS		Weight-for-height SDS	
		N	Mean $\pm$ SD	N	Mean $\pm$ SD	N	Mean $\pm$ SD
0-4 months	f	5	- 1.84 $\pm$ 1.73	5	- 2.84 $\pm$ 1.17	5	- 1.02 $\pm$ 0.80
	m	9	- 2.88 $\pm$ 1.67	9	- 3.28 $\pm$ 1.37	9	- 1.11 $\pm$ 0.76
18-24 months	f	7	- 0.45 $\pm$ 0.82	7	- 0.95 $\pm$ 1.26	7	- 0.47 $\pm$ 1.06
	m	10	- 0.23 $\pm$ 0.99	10	- 0.32 $\pm$ 0.70	10	- 0.21 $\pm$ 0.72
60-66 months	f	6	- 1.48 $\pm$ 0.97	6	- 1.19 $\pm$ 0.95	6	- 0.53 $\pm$ 0.47
	m	8	- 1.50 $\pm$ 0.54	8	- 1.79 $\pm$ 0.84	8	- 1.16 $\pm$ 1.09
102-108 months	f	6	- 1.61 $\pm$ 0.81	6	- 1.49 $\pm$ 0.54	6	- 0.61 $\pm$ 0.42
	m	4	- 0.65 $\pm$ 0.47	4	- 0.48 $\pm$ 1.00	4	0.40 $\pm$ 1.71

FIG 3.1. A-C. Mean and 95% confidence intervals for height SDS (a), weight SDS (b) and weight-for-height SDS (c) in CF patients from 0 up to 120 months.

FIG 3.2. A-C. Correlations between catch-up growth for height (a), weight (b) and weight-for-height (c) with severity of initial growth failure expressed as worst initial SDS for height, weight and weight-for-height.

female and male CF patients (-1.84 and -2.88 respectively). Beyond the age of 2 years the mean height SDS was nearly normal (-0.45 and -0.23 respectively). Weight SDS followed the same pattern with even worse initial SDS. The slight height and weight SDS differences between boys and girls were not significant. Female and male patients showed similar initial weight-for-height SDS (-1.02 and -1.11 respectively). The mean weight-for-height SDS of male patients increased up to 0.40 in the 102-108 months period and was by that even better than in normal Dutch boys. While the mean weight-for-height SDS (-0.61) of female patients remained slightly, but not significantly, below mean values for normal girls.

figure 3.1a-c shows the mean and 95% confidence intervals for height, weight and weight-for-height SDS for boys and girls taken together during the study period. The height SDS started at a very low value, increased to the normal range at the age of two years, but subsequently decreased again, reaching a plateau of -1.5 SD at the age of 5-10 years. Weight SDS had an identical pattern with a plateau of -1 SD at the age of 3-10 years. Initially weight-for-height was much less decreased and, besides a short period around 4-5 years, it was from 1.5 years on in the normal range compared to Dutch children.

A strong positive correlation was found between catch-up growth and the severity of initial growth failure in our patients. Correlation coefficients (r) for height, weight and weight-for-height were resp. -0.86 ($P<0.001$), -0.80 ($P<0.001$) and -0.60 ($P<0.05$) (figure 3.2). While catch-up growth for weight and height were significantly related to each other ($r=0.78$, $P<0.01$).

3.5. DISCUSSION

Also in our small but longitudinal study, at time of diagnosis (before 4 months of age in all our patients) we demonstrated, in both boys and girls with CF, an obvious decreased height and weight SDS, while weight for height SDS was only slightly decreased. In our study mean height, weight and weight-for-height SDS normalised at the age of 18-24 months, and subsequently decreased somewhat at the age of about five years. Previous studies in CF patients have shown subnormal initial height and weight SDS [1-7, 11-13]. Compared to the latter reported data, our patients showed a more severe initial growth retardation. This could partly be due to the fact that there is no standard screening-program for CF in the Netherlands. On the other hand diagnosis was clear before the age of four months in all of our patients. Important to notice is the almost complete catch-up growth in the first 2 years after diagnosis, which implicates that this growth retardation was reversible.

Karlberg et al. [4] evaluated linear growth in children with CF from birth to 8 years of age. Height SDS were reported to be subnormal up to one year of age, improved up to a height of -0.3 SD at five years of age and even became positive between 5 and 8 years of age. These findings only differ from ours, concerning the partly reverse of catch-up growth found in our patients at about 5-10 years. In a cross-sectional British study, reporting cross sectional growth of 3056 children with CF, showed also a plateau in

growth, between 1 and 10 years. The plateau was approximately 0.5 SD below the population mean. Before the age of one, both weight and height were below those of the general population, normalized by two years of age and did not change thereafter. After the age of 10 years the SDS declined [12]. The improvement after the age of one year was partly explained by inclusion of late diagnosed mild cases in older cohorts and partly reflected catch-up growth after effective treatment. These cross-sectional study results do perfectly agree with our own longitudinal data. Because of the early diagnosis in all our cases we excluded the possible bias due to the inclusion of later diagnosed mild cases.

Our results show that in CF patients, catch-up growth was strongly related with the severity of the initial growth failure for both weight and height. For weight-for-height SDS this relationship was less impressive: in a previous study reporting catch-up growth of children with congenital heart disease (VSD, Fallot), we showed in severely growth retarded patients only catch-up growth for height to be linearly related to the severity of the initial growth retardation [17].

In the present study, catch-up growth for weight and height were strongly related to each other. The latter finding could be expected as weight and height are interrelated and it also can be explained as a regression to the mean.

As expected weight-for-height is only slightly abnormal pointing to the presence of stunting more than wasting in our patients. Stunting is defined as height-for-age two or more standard deviations below the median and implies chronic malnutrition. Wasting is defined as a low weight-for-height and implies acute malnutrition [15,18,19]. Especially the “catch-down” period found in CF patients of 5–10 years old, consists mainly of “stunting” as weight-for-height remained perfectly normal in most patients.

In conclusion, catch-up growth for weight, height and weight-for-height in our CF patients occurs in the first years after diagnosis and results in normalisation of SDS for each parameter around the age of two years. A slight decrease in both weight and height SDS occurs later on while weight-for-height remains normal.

Our results show “wasting” to be a rare event in the evaluation of these patients growth disturbances. Consequently, both height-for-age and weight-for-age are more appropriate than weight-for-height growth charts for the nutritional follow-up of this chronic disease.

Furthermore we think that these findings support the use of a screening program for CF, in order to prevent stunting at an early and sensitive age period.

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CHAPTER FOUR

Potential short and long term effects of lansoprazole on body composition in children with Cystic Fibrosis

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Submitted

4.1. ABSTRACT

Background: In a previous, short-term, study on the additional effect of a proton pump inhibitor (PPI) in Cystic Fibrosis (CF) patients, we were able to demonstrate an improvement in fat absorption and total body fat (TBF), but not fat free mass (FFM).

Objective: The present study was performed to evaluate both the short-term (1 yr) and long-term (3 yrs) effects of adjuvant therapy with a PPI on body composition in CF.

Design: In this prospective, observational study 14 CF patients with pancreas insufficiency used lansoprazole, 15 mg daily for 3 years, as adjuvant treatment. Body composition was assessed at the start of the study and at 6, 12 and 36 months by dual energy X-ray absorptiometry (DEXA). Results were expressed as SD scores (SDS), using a large group of normal Dutch children. Non-parametric tests (Wilcoxon) were used.

Results: The decreased TBF, bone mineral content (BMC) and bone mineral density (BMD) SDS increased significantly (P values: 0.006, 0.022, and 0.013 respectively) at one year, while FFM did not change ($P=0.551$). After 3 years all three parameters regressed and were not different from starting values anymore.

Conclusion: The addition of a PPI to the standard treatment of young CF patients showed an improvement of TBF, BMC and BMD at one year, whereas its levels after 3 years returned to starting values. As this study was not controlled, it only suggests that, at least temporarily, besides TBF also BMC and BMD might improve during adjuvant therapy with PPIs in CF. Furthermore we advocate that regular assessment of body composition by DEXA in CF allows a better monitoring of the nutritional condition.

4.2. INTRODUCTION

Cystic Fibrosis (CF) is an autosomal recessive inherited disorder, characterized by a defect in the chloride and bicarbonate transport regulating channel at the luminal side of exocrine cells, the cystic fibrosis transmembranous conductance regulator protein (CFTR). One of the most serious consequences of this defect is pancreatic insufficiency, which induces mainly fat malabsorption. This pancreatic insufficiency is only partially corrected by exogenous pancreatic enzymes [1]. In two, separate, studies we have shown that lansoprazole, a proton pump inhibitor (PPI), used as adjuvant therapy, significantly improved both steatorrhea and the nutritional status of CF patients after 3 months [2] and one year [3] use of lansoprazole, a PPI. At the same time lung hyperinflation decreased in those CF patients [3]. The purpose of a PPI is to improve fat absorption by decreasing gastric acid production resulting in a lower acid duodenal pH [4-6]. This is necessary because of decreased pancreatic ductular bicarbonate secretion, resulting in a low duodenal buffering capacity and a low luminal pH of the intestine.

Malabsorption in CF enhances the poor nutritional status, while a relationship between nutritional status, growth and lung function has been demonstrated in these patients [7-12]. The present study was performed primarily to evaluate short-term (0-1 year) and long-term (3 years) effects of adjuvant therapy with PPIs on body composition (TBF, FFM, BMC and BMD) as measured by DEXA in young CF patients treated in a standard way according to the Dutch "CBO consensus" [13].

The Medical Ethical Committee of the University Hospital of Maastricht approved the study protocol and informed consent from the parents and from children older than 11 years of age was obtained before onset of the study.

4.3. PATIENTS AND METHODS

PATIENTS

14 CF patients (9 males and 5 females), with pancreatic insufficiency and mild lung disease, were included in a prospective study. In all patients the diagnosis of CF was confirmed by sweat testing (Chloride more than 60 mmol/l) and all demonstrated persistent steatorrhea despite regular pancreatic enzyme replacement therapy. Except for one patient, the oldest one, all subjects were studied during 3 years of adjuvant therapy with 15 mg lansoprazole daily. Diet and exogenous pancreatic enzymes were adjusted before the study, and checked by our dietary consultant. All other therapy was standardized, according to the Dutch "CBO consensus" [13]. As this study was an open study, the patients were used as their own control, and changes were expressed as changes in standard deviation scores (SDS). Therefore anthropometric data of our patients (height, weight and BMI) were expressed as SDS using a large healthy Dutch reference population [14,15]. To include only stable patients, inclusion criteria for this study were no recent infection or exacerbation, and a forced expiratory volume in 1 second (FEV₁) of more than 70% of predicted in the preceding

TABLE 4.1. Patient characteristics at start (N=14)

Item		Mean	Median	SD	Range
Age	(yr)	9.74	8.60	3.38	4.6 / 17.2
Height	(cm)	133.9	131.5	16.6	107.0 / 162.3
	(SDS)	-1.07	-1.11	0.59	-2.17 / 0.00
Weight	(kg)	27.7	25.6	9.9	15.5 / 52.3
	(SDS)	-1.27	-1.44	0.74	-2.39 / 0.27
Acid Steatocrit	(%)	33.3	32.7	17.7	5.4 / 65.7
FEV ₁	(% predicted)	83.9	85.0	17.2	58 / 112
FVC	(% predicted)	86.6	87.0	13.0	64 / 109

SDS = standard deviation score

year. But the most important inclusion criterion was a mean pre-study stool acid steatocrit (an index of steatorrhea) of more than 25% (normal <20%), despite an intake of at least 2000 IU lipase/kg body weight/day (mean use 4020 IU/kg/day). Patient characteristics at inclusion are summarized in table 4.1.

FECAL FAT CONTENT

steatorrhea was monitored by measuring fecal acid steatocrit as previously described [16,17]. Here results of fecal fat content are expressed as percentage of fat layer of the addition sum of fat and solid layer. Fecal acid steatocrit has been shown to correlate well with chemical analyses of fecal fat ($r = 0.83$)[18]. We chose this method in order to repetitively and easily monitor fecal fat content, which was measured every 2 weeks during the first year of PPI therapy and at 3 years.

BODY COMPOSITION PARAMETERS

Body composition was assessed by dual energy X-ray absorptiometry (DEXA). This non-invasive method allows an easy and rapid (20 minutes) evaluation of body composition. During one assessment, total body fat (TBF), fat free mass (FFM), bone mineral content (BMC) and bone mineral density (BMD) are measured. A software pediatric program (Pediatric Beta Software: Lunar 1.5) was used. The radiation dose of this examination is very low (<5 mRem) when compared with the standard background radiation exposure (82 mRem/year), and is therefore considered to be the method of choice to measure body composition in young children. Results were compared with those of a large ($n=403$) healthy Dutch reference group matched for age and sex and expressed as SDS [19]. In the latter study the variation in TBF was 4%, which is twice the variation of adults. DEXA results were evaluated at start, after 6, 12 and 36 months use of a PPI.

STATISTICS

Non-parametric tests (Mann-Whitney, Wilcoxon) were used to compare SDS of our patient group with those of the reference populations at start and at different intervals. By using a Non-parametric test, we avoided making an assumption of normality, which cannot be tested in only 14 subjects. Differences with respect to baseline were considered significant when the two-sided P -value was 0.05 or less. The statistical package SPSS (version 10.0 for Windows, 1999, SPSS Inc., Microsoft, Chicago, IL) was used for data management and analysis.

4.4. RESULTS

FECAL FAT CONTENT

Shortly after the addition of lansoprazole a significant decrease in fecal fat content (acid steatocrit) was noticed. As a group, mean acid steatocrit decreased from 33.3% (± 17.8) at the start to 23.7% (± 11.0) at 3 months ($P < 0.01$) and remained at that level thereafter. The mean value during the first year of treatment was 24.0% (± 9.8 ; $P < 0.05$). After use of a PPI for 3 years, mean acid steatocrit was still 23.4% (± 11.8).

ANTHROPOMETRY

Box plots for height and weight SDS during 3 years treatment with a PPI are shown in figure 4.1. The mean (SD) bodyweight SDS changed slightly but not significantly from -1.27 (± 0.74) before to -1.12 (± 0.82) at 1 year and -1.28 (± 0.95) at 3 years. Mean (SD) height SDS hardly changed from -1.07 (± 0.59) before the study to -1.02 (± 0.61) at 1 year and -1.05 (± 0.82) at 3 years all NS. Table 4.2. shows the mean and median values of the BMI, in girls and boys separately. Mean (SD) BMI at the start of the study was somewhat decreased compared to values of normal children. During the first year it showed an increase of 0.5 kg/m^2 or 0.15 SDS, which is exactly what could be expected in healthy children [20]. At 3 years, however, hardly any further increase of BMI was assessed, whereas a further increase of 1.2 kg/m^2 is normal in this age group, resulting in SDS values comparable with starting values. Remarkable is the discrepancy between both sexes, favouring boys.

BODY COMPOSITION

Means (SD) and medians (range) for body composition data are shown in table 4.3. and in figure 4.2. After using a PPI for 6 months a significant increase in SDS of TBF was found ($P = 0.035$). It is worth mentioning that TBF in grams increased by 32.6% and, although a part of this increase will be normal growth, it is far beyond the 4% of variation in the measurement. After 1 year SDS of TBF, BMC and BMD all increased significantly, ($P = 0.006$, 0.022, and 0.013 respectively), while SDS of FFM did not change. All parameters at 3 years decreased and were not significantly different from starting values.

TABLE 4.2. Mean and median Body Mass Index during 3 years treatment with lansoprazole, in kg/m^2 and SDS, of the 14 CF patients, with separate numbers for both sexes.

Sex		At start		At 6 months		At 1 year		At 3 years	
		kg/m^2	SDS	kg/m^2	SDS	kg/m^2	SDS	kg/m^2	SDS
F	Mean	15.86	-1.24	15.95	-1.29	16.37	-1.21	15.89	-1.38
	SD	2.40	0.64	2.32	0.60	2.49	0.61	1.62	0.71
	Median	14.55	-1.21	15.37	-1.21	15.37	-1.24	15.23	-1.36
	Range*	5.64	1.61	5.84	1.64	6.09	1.58	3.44	1.90
M	Mean	14.45	-1.00	14.96	-0.66	14.90	-0.77	15.39	-0.90
	SD	1.29	1.05	1.75	1.33	1.59	1.20	2.17	1.35
	Median	14.13	-1.22	14.44	-1.05	14.47	-0.81	14.73	-1.45
	Range	3.86	3.30	5.18	4.24	4.90	3.90	6.76	4.42
Total	Mean	14.95	-1.09	15.32	-0.88	15.43	-0.93	15.54	-1.08
	SD	1.81	0.91	1.95	1.14	1.99	1.03	1.97	1.15
	Median	14.31	-1.21	14.48	-1.16	14.71	-0.92	14.86	-1.41
	Range	7.25	3.30	6.89	4.25	7.19	3.90	6.76	4.48

*Range is given in one number (highest – lowest value).

FIGURE 4.1. Box plots (medians and 95%) of height and weight SDS, during 3 years treatment with lansoprazole in CF patients.

TABLE 4.3. Mean (SD) and Median (Range) of SDS DEXA body composition parameters, before and during 3 years use of lansoprazole in CF patients.

SD scores		At start (n=14)	At 6 months (n=14)	At 1 year (n=14)	At 3 years (n=13)
Age	Mean (SD)	9.7 (3.4)	10.2 (3.4)	10.7 (3.4)	12.3 (2.8)
	Median (range)	8.6 (12.6)	9.1 (12.6)	9.6 (12.6)	11.9 (9.6)
FFM	Mean (SD)	-1.82 (0.67)	-1.90 (0.78)	-1.78 (0.63)	-2.33 (0.83)
	Median (range)	-1.93 (2.54)	-1.93 (3.31)	-1.83 (2.25)	-2.52 (2.61)
TBF	Mean (SD)	-0.97 (0.49)	-0.73 (0.73)*	-0.68 (0.70)*	-1.03 (0.74)
	Median (range)	-0.95 (1.73)	-0.88 (2.57)*	-0.85 (2.35)*	-1.32 (2.82)
BMC	Mean (SD)	-1.78 (0.76)	-1.71 (0.82)	-1.55 (0.85)*	-1.80 (1.01)
	Median (range)	-2.03 (2.35)	-1.89 (2.67)	-1.72 (2.65)*	-1.62 (3.14)
BMD	Mean (SD)	-0.99 (1.12)	-0.95 (1.17)	-0.67 (1.18)**	-0.91 (1.48)
	Median (range)	-0.88 (3.55)	-0.92 (4.18)	-0.65 (3.71)**	-0.92 (4.54)

SDS standard deviation score (= Z score)

Range is given in one number (highest – lowest value).

All interval numbers (except age) are compared with starting values, by Wilcoxon rank sum test; 2 sided, 2 related: * P value < 0.05, ** P value < 0.01

FIGURE 4.2. Changes in SDS for body composition parameters by DEXA during 3 years treatment with lansoprazole in CF patients.

4.5. DISCUSSION

In this observational study, the addition of a PPI demonstrated effects, which were not all expected. The short-term (1 year) effects on fecal fat content (acid steatocrit), and body composition (BMI, TBF, BMC and BMD) suggest a beneficial effect, but on the long-term (3 years) this was not confirmed and values dropped back to starting values. Furthermore no effect on body height, weight and FFM was shown. Although malnutrition is common knowledge in CF, we wondered why the starting values of body height and weight were decreased so much, and more than expected. First, our CF population is from the southern part of the Netherlands, where adults, like children, have been shown to be at least 5 cm shorter than in the northern part where our reference growth data come from. Second, CF patients are known to have a delayed maturation and, as a mean, enter puberty with a delay of about 1 year [21]. Both explanations will result in an overestimation of our patients' growth deficit, and therefore in an underestimation of the body composition parameters.

A decrease in fecal fat content during the use of a PPI is not unknown: besides our earlier reports in children [2,3], it has also been demonstrated previously by Heijerman et al in adults [22]. Therefore the demonstrated favourable and persistent effect of the use of PPIs on fecal fat content, as measured by the acid steatocrit, was expected. The demonstration of an improvement in body fat and bone mass during the first year with the apparent loss at 3 years is much more difficult to interpret, as this is not a controlled study. A placebo effect could also be responsible for the observed changes. However, we could demonstrate the same changes in body composition in two separate studies. The natural course of CF disease shows a decrease in weight and peak bone mass during the second decade [23-27]. Recently, in a 3-year longitudinal study, Stettler et al. [11] showed growth of boys 5-10 years old to be impaired not only for height and weight, but also for FFM and TBF. For girls in that age group these changes are less striking. Therefore during the 3 years of follow up we might have expected a continued decrease in body weight and height, and body composition parameters like BMI, TBF, FFM, BMC and BMD. As we could not demonstrate this expected decrease in 3 years in our patients whose body composition data were similar to their starting values, the addition of a PPI might suggest a delay in the expected decay of weight and bone acquisition over 3 years. According our dietician during the addition of a PPI either the diet and the use of pancreatic enzymes did not change significantly [28]. Furthermore we could demonstrate an improvement in skinfold measurements too. Both the suprailiacal skinfold, and the sum of four skinfolds improved significantly ($P < 0.05$) and correlated very well with TBF as measured by DEXA ($r^2 = 0.86$) [29]. As we have to interpret all these results cautiously, we therefore only suggest that the use of PPIs in young CF patients might have some effects on body composition.

More remarkably is the improvement in bone mineral status during PPI treatment, which was unexpected. A high prevalence of bone disorders, such as osteoporosis with an increased risk for bone fractures, has been reported in CF patients. Several explanations have been suggested such as malnutrition, especially vitamin D and calcium malabsorption, but also relative inactivity, hormonal alterations and increasing age of

CF patients. Bone metabolism is not static but a dynamic process, being the result of bone formation and resorption. Evaluation of bone turnover markers in CF patients, have shown an imbalance between bone formation and resorption [30]. Bone mineralization in CF patients has been related to age, pubertal stage, body mass, energy expenditure, illness severity, glucocorticoid therapy and gonadal function. Changes in each of these factors because of the disease can therefore cause a poor bone mineral status [27,31]. Previous studies have shown that serum parathyroid hormone and calcium, carbohydrate intake and activity had limited predictive value, while daily calcium intake and CFTR-genotype had no predictive value on bone mineral status. In many studies a correlation between bone mineral content and body composition, in particular fat-free mass has been demonstrated. However, it might be too simple to explain the defective bone mineral status by either malnutrition or malabsorption since it was also found in patients with completely normal stature and even in pancreatic sufficient patients. In the present study we demonstrated in young pancreas insufficient CF patients, evaluated by DEXA, a significant improvement in BMC and BMD, during adjuvant therapy with a PPI. As height SDS did not change, the increase in BMD might be real and not due to improved growth. The response was not immediate, but occurred between the sixth and the twelfth month of treatment, while malabsorption improved within 3 months. The delay in effect on BMC and BMD explainable by seeing bone turnover as a cyclic process, lasting for about 120 days. The first 20 days there is a bone resorption phase, followed by bone formation for some 100 days [32]. Therefore it is likely that more time is needed to reflect improvements in bone mineral status, and one cycle might be too short to show significant changes. Improvement of BMC and BMD point to a potential beneficial effect of PPI on bone formation. Possible explanations are increased absorption of minerals and fat soluble vitamins, like vitamin D and K. Particularly vitamin K deficiency, a co-factor in the carboxylation of osteocalcin (OC) a marker of bone formation, might play a role in the decreased bone mineral status in CF patients. In order to elucidate the relation between vitamin K and bone turnover markers we performed a study in our department. Preliminary results point to such an under-carboxylation of OC in CF (to be published separately).

In conclusion, adjuvant therapy with a PPI in young CF patients revealed that potential short term (1 year) improvements returned to starting values in the long term (3 years). The effect of PPIs on fecal fat content seems not to be temporary, as acid steatocrit values at 3 years were still as low as at 1 year. It might be that comparison of CF patients, having a delayed puberty, with a healthy population resulted in an apparent decrease of body composition SDS at 3 years compared to pre-pubertal starting values. Another possibility might be that the natural course of CF in 3 years of follow-up is delayed somewhat. Therefore one might conclude that PPIs have a positive, undetected, long-term effect on body composition, but due to its design this statement cannot be proven by the present study. To elucidate further the role of PPIs in improving body composition and bone mineral abnormalities in CF, more placebo-controlled studies are necessary.

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CHAPTER FIVE

Changes in pulmonary hyperinflation and bronchial hyperresponsiveness following treatment with lansoprazole in children with Cystic Fibrosis

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5.1. ABSTRACT

Background: In this prospective, preliminary, open study in 14 children with Cystic Fibrosis (CF), we evaluated the effect of one year adjuvant therapy with lansoprazole, a proton pump inhibitor (PPI), on growth, fecal fat losses, body composition and lung function parameters.

Methods: Only stable patients with pancreas insufficiency were included and all their data were compared to those of large Dutch pediatric normal reference populations.

Results: During the use of a PPI mean weight and height did not change significantly, while body mass index improved significantly ($P<0.05$). An immediate significant and persistent reduction of fecal acid steatocrit ($P<0.05$) was demonstrated. Compared to normal Dutch children the CF patients showed significantly decreased SD-scores (SDS) for total body fat (TBF; -0.966) and fat free mass (FFM; -1.826). Under lansoprazole, TBF improved significantly ($P<0.05$) while mean FFM remained unchanged. Meanwhile a significant improvement of total lung capacity (TLC; $P<0.05$), residual volume (RV; $P=0.055$), and maximal inspiratory mouth pressure (PI_{max}; $P=0.002$) was demonstrated. These findings confirm the presence of hyperinflation in CF, which seems to decrease during the use of a PPI. Daily recordings of peak expiratory flow (PEF) showed a maximal diurnal variability of 28% of recent best PEF and minimal morning PEF of 72% of recent best PEF, which confirms that bronchial hyperresponsiveness (BHR) is increased in CF.

Conclusion: We conclude that adjuvant therapy with lansoprazole in our young CF patients with persistent fat malabsorption, decreased fat losses and improved total body fat. Meanwhile lung hyperinflation decreased, which can partly explain the inspiratory muscle performance improvement. The coincidence between improvement of bodycomposition and lung hyperinflation might suggest a relationship between these two parameters. Further research is necessary in order to confirm a relationship and to elucidate the mechanisms involved.

5.2. INTRODUCTION

CF is the most common lethal hereditary disease amongst Caucasians. The CF gene is located on chromosome 7 and so far more than 900 mutations of this single gene have been reported. The basic defect is a dysregulation of the chloride channels resulting in dehydration of the luminal surface of exocrine cells. In the lungs this leads to an increase of mucus viscosity, inhibiting normal ciliary clearance. Bacterial colonization, infection and chronic pulmonary inflammation develop subsequently. Pulmonary complications account for most of the morbidity and mortality in CF patients. Besides the respiratory complications, gastroenterological abnormalities give rise to significant morbidity. In children with CF, fat malabsorption leads to malnutrition and growth failure, often associated with increased catabolism. Exogenous administration of pancreatic enzymes improves, but does not reverse fat malabsorption completely. It has been hypothesized that the use of a PPI could improve the effectiveness of the administered pancreatic enzymes. The aim of adding these drugs is to decrease gastric acid production. This is necessary in CF because of a decreased buffering capacity of the pancreatic juices, due to decreased or absent bicarbonate. If duodenal pH is less than 5 early dissolving of the enteric coating of the administered enzyme capsules is hindered, because this only takes place at a duodenal pH above 5.5. [1,2] In addition this low pH may also inactivate pancreatic enzymes. Both processes result in suboptimal functioning of these enzymes, accompanied by incomplete absorption of nutrients like fat.

A favourable effect of PPI on fat absorption in adult CF patients was demonstrated by Heijerman et al. [3] We confirmed this favourable effect in a pilot study in children [4]. In the latter short-term (3 months) study, the use of lansoprazole led to a significant increase of TBF or fat mass while FFM or lean body mass did not improve. Although it has frequently been suggested that changes in body composition can affect lung function and especially respiratory mechanics, this has, until now, hardly been proven in CF patients. Research to support this hypothesis is sparse. Steinkamp and von der Hardt [5] demonstrated that nocturnal tube feedings improved nutritional status and lung function in malnourished CF patients. More recently Ionescu et al. [6] demonstrated that in the last stages of CF, survival was negatively associated with a BMI < 20 kg/m² and low FFM, while sustained maximum inspiratory pressure seemed to be more impaired than peak maximum inspiratory pressure.

By impairing lung growth, suboptimal increase of body cell mass has also been suggested to be associated with progressive lung dysfunction in CF [7].

Apart from its effect on fat absorption and body composition, a PPI could have a favourable influence on BHR by decreasing gastroesophageal reflux (GER), possibly justifying its use in CF patients.

5.3. AIMS OF THE STUDY

The present open study was performed in 14 children with stable CF, and primarily designed in order to investigate the effect of long term addition of a PPI on body composition. Secondly the relation between changes in body composition and changes in lung function of dynamic and static lung function as well as respiratory muscle parameters, were examined. Thirdly we evaluated the effect of decreased acid secretion and possibly decreased GER on BHR.

5.4. MATERIALS AND METHODS

PATIENTS

14 stable CF patients (9 males) with pancreatic insufficiency and mild lung disease were studied before and during one year of treatment with 15 mg lansoprazole daily. Diet and exogenous pancreatic enzymes did not change during the intervention. We conducted an open study for practical reasons and because we preferred to use our patients as their own control, as every patient is unique and not exactly comparable to any other CF patient. We therefore compared them to large normal (Dutch) populations and expressed changes as changes of SD-scores. In order to reduce a possible placebo effect only stable patients were included in the study, demonstrated by the high pre-study FEV₁, which was not different from the baseline FEV₁. Criteria for inclusion in the study were a mean prestudy stool acid steatocrit (an index of fat absorption) of more than 25% (normal <20%), despite an intake of at least 2000 IU lipase/kg bodyweight (mean use 4020 IU/kg). Further our CF patients had to be able to perform reliable lung function tests and in the prestudy year their FEV₁ had to be above 70% of predicted (mean prestudy year FEV₁ was 85.2 ±13.6% of predicted). Finally they had to be able to use a Peakflow meter and fill-in diary cards correctly. Verbal and written informed consent was obtained before onset of the study and the study

TABLE 5.1. Patient characteristics at start

Item		Mean	SD	Range
Age	(yr)	9.74	3.38	4.6 / 17.2
Height	(cm)	133.9	16.6	107.0 / 162.3
	(SDS)	-1.068	0.590	-2.166 / 0.003
Weight	(kg)	27.7	9.9	15.5 / 52.3
	(SDS)	-1.270	0.739	-2.393 / 0.272
Acid Steatocrit	(%)	33.3	17.7	5.4 / 65.7
FEV ₁	(% predicted)	83.9	17.2	58 / 112
FVC	(% predicted)	86.6	13.0	64 / 109

For abbreviations see list of abbreviations (page 6).

protocol was approved by the medical ethical committee of the university hospital of Maastricht. Patient characteristics at baseline are given in Table 5.1.

ANTHROPOMETRY

Anthropometric measurements were performed in triple by one single investigator using the same instruments, a Harpenden stadiometer (Holtain limited, Crymych, D Y Fed., UK) and an electronic weight scale (Seca delta, model 707). Results are expressed as SDS using Dutch reference values collected by Gerver and de Bruin [8,9]. The subjects were all measured barefoot and dressed in underwear.

FECAL FAT CONTENT

steatorrhea was monitored by measuring the fecal acid steatocrit as previously described [10,11], where results are expressed as fat / fat+solid layer in %. Fecal acid steatocrit has been shown to correlate well with chemical analyses of fecal fat ($r = 0.83$) [12]. We chose for this method in order to repetitively and easily monitor fecal fat content during the study.

BODY COMPOSITION

Body composition was assessed by DEXA [13]. This non-invasive method allows an easy and rapid (20 minutes) evaluation of body composition. During one assessment TBF, FFM, BMC and TBD are measured. A software pediatric programme (Paediatric Beta Software: Lunar 1.5) was used. The radiation dose of this examination is very low (<5 mRem) when compared with the standard background radiation exposure (82 mRem/year), and is therefore considered to be the method of choice to measure body composition in young children. Results were compared with those of a large ($n=403$) age matched normal Dutch reference group [14] and expressed as SDS.

LUNG FUNCTION

Lung function tests were performed by an experienced lung function technician, using a Jaeger Masterlab pneumotachograph and bodybox. Calibration of the apparatus was performed twice daily. Only lung function data with FVC variability of less than 5%, on 3 consecutive manoeuvres, were accepted for data management. Lung function parameters were expressed as % of predicted (= mean of normal children, for height and sex) according to the Zapletal reference values [15]. Maximal mouth pressures were measured at RV for inspiratory pressure (PI_{max}) and at TLC for expiratory pressure (PE_{max}) according to the method of Black and Hyatt [16].

BRONCHIAL HYPERRESPONSIVENESS

PEF was measured at home, using a "Personal Best" PEF meter (Respironics HealthScan Products, Europe, Belgium) four times daily. Instruction was given about the correct technique and reading of the PEF meter. Three consecutive manoeuvres were carried out and recorded, but only the best of these 3 was used for data management (also in the so-called minimal morning PEF!). In order to measure changes in

BHR several PEF characteristics were calculated for each weekly recordings every month [17]:

- ◆ Mean diurnal variability (in L/min).
- ◆ Maximal diurnal variability: the difference between the worst morning and best evening PEF of that week (expressed as % of recent (that week) best PEF).
- ◆ Minimal morning PEF (expressed as % of recent best PEF).
- ◆ The difference between the best morning and the best evening PEF (in L/min).

RESTING ENERGY EXPENDITURE (REE) AND DIET

Basal metabolic rate or REE was measured by indirect calorimetry [18] (ventilated hood), and compared to Schofield's age related normal values for children [19]. Dietary evaluation was carried out using an extensive booklet for three days recording of the diet of these CF patients. From the recorded data lipase intake (FIP-E), dietary fat (gram/day), protein (gram/day) and energy (KJ/day) content were calculated.

EXPERIMENTAL PROTOCOL

The following measurements were obtained over a period of one year, according to the protocol. Before start and at each 3 monthly visit, anthropometric data were obtained (5x). Three faecal samples before start of the study and two samples were taken monthly during the 12 months study period, providing 27 specimens for analysis of acid steatocrit.

Totalbody DEXA was performed every 6 months (3x). Lung function tests: Flow Volume curve, Bodybox and PI/ PE-max were performed at each visit (5x) and REE was measured every 6 months (3x). Diary cards with PEF were recorded during one week, monthly (13x). The dietary intake was recorded before start, at 6 and 12 months (3x).

STATISTICS

Data are presented in tables as means and standard deviations, while in the figures medians and quartiles are presented as changes from baseline. We tested the response to the treatment for all measured parameters. To maximize the power of the tests, we compared the mean of the whole intervention period to the pre-trial (baseline) values. In that way, the power of the test is optimized, since measurement errors and within-subject variation tends to cancel out. A second advantage of this method is that we avoided doing 4 different tests for essentially the same hypothesis, which would inflate the type-I error probability beyond the nominal 5% [20]. The price to pay is superficially a less detailed picture of trends within the trial period. However, with a small trial as the present one, we would not have any real power to detect such trends. If a clear trend is visible we noted them in the results and the figures show some of these trends. By using the non-parametric Wilcoxon signed ranks test, we furthermore avoided to make the assumption of normality, which is untestable with only 14 subjects. All data are summarized in the Tables 5.2-4. Differences with respect to baseline were considered significant when the two-sided *P*-value

was 0.05 or less. The statistical package SPSS (version 7.5 for Windows, 1996) was used for data management and analysis.

5.5. RESULTS

ANTHROPOMETRY

Mean ($\pm$ SD) bodyweight SDS increased from -1.270 (± 0.739) to -1.085 (± 0.827) at 9 months ($P < 0.05$). However the mean value during the study was -1.118 (± 0.817 ; NS). Mean height SDS showed a slight, but not significant, increase from -1.068 (± 0.590) before to -1.017 (± 0.608) mean during the study. BMI was considerably decreased before the study 14.95 (± 1.81 kg/m²) but improved significantly to 15.38 (± 1.92 kg/m²; $P < 0.05$) (Table 5.2).

FECAL FAT CONTENT

An immediate and significant decrease in fecal fat content (acid steatocrit) was noticed, shortly after the addition of lansoprazole. The mean acid steatocrit decreased from 33.3% ($\pm 17.8\%$) at the start to 23.7% ($\pm 11.0\%$) at 3 months ($P < 0.01$) and remained at that level thereafter, the mean value during treatment was 24.0% ($\pm 9.8\%$; $P < 0.05$).

BODY COMPOSITION

Body fat assessed by DEXA performed at 6 months showed a significant improvement, persisting during the entire study. Mean TBF SDS had increased from -0.966 (± 0.489) before to -0.707 (± 0.695) during PPI ($P < 0.05$). In contrast mean FFM SDS did not change, from -1.820 (± 0.673) before to 1.843 (± 0.687 ; NS) during PPI (Table 5.2 and Figure 5.1).

LUNG FUNCTION

Mean TLC decreased from 109.5 ($\pm 16.6\%$ of predicted) before to 100.3 ($\pm 7.3\%$; $P < 0.05$) during PPI, mean RV showed a nearly significant decrease from 179.6 ($\pm 73.1\%$) before to 141.5 ($\pm 29.9\%$; $P = 0.055$) during PPI, mean ITGV decreased from 128.0 ($\pm 42.9\%$) before to 111.0 ($\pm 13.1\%$; NS) during PPI and mean ERV increased from 71.9 ($\pm 15.6\%$) to 78.6 ($\pm 20.6\%$; NS) during PPI. When ratio's of these static lung volumes were calculated, no significant change was noted, the RV/TLC ratio changed from 40.1 ($\pm 10.1\%$) before to 35.2 ($\pm 7.2\%$) during PPI, while the ITGV/TLC ratio changed from 55.7 ($\pm 9.2\%$) before to 54.0 ($\pm 4.6\%$) during PPI (Table 5.3). Dynamic lungfunction parameters did not change significantly. Mean FEV₁ changed from 83.9 ($\pm 17.2\%$) to 81.8 ($\pm 16.9\%$; NS) during PPI, mean FVC from 86.6 ($\pm 13.0\%$) to 87.6 ($\pm 13.1\%$; NS) during PPI. Mean sGAW did not change, from 0.864 1/kPa.s (± 0.257) to 0.854 (± 0.219 ; NS). Also mean pneumotachographic PEF did not change, from 90.6 ($\pm 18.7\%$) to 87.9 ($\pm 15.4\%$; NS) during PPI (Table 5.3).

TABLE 5.2. Mean values and statistics of anthropometry, REE, body composition and diet parameters before and during PPI.

Parameter	Mean (SD)	Mean (SD)	Wilcoxon <i>P</i> =	Number
	Before PPI	During PPI		
Weight (SDS)	-1.270 (0.739)	-1.118 (0.817)	0.177	14
Height (SDS)	-1.068 (0.590)	-1.017 (0.608)	0.198	14
BMI (kg/m2)	14.95 (1.81)	15.38 (1.92)	0.035	14
Acid Steatocrit (%)	33.3 (17.8)	24.0 (9.8)	0.030	14
TBF (SDS)	-0.966 (0.489)	-0.707 (0.695)	0.011	14
FFM (SDS)	-1.820 (0.673)	-1.843 (0.687)	0.826	14
REE (% pred.)	111.7 (12.1)	110.7 (9.1)	0.778	14
Lipase intake (IU/kg)	4020 (970)	3580 (880)	0.198	14
Fat intake (gr/d)	90.5 (18.8)	81.9 (17.0)	0.056	14
Energy intake (kJ/d)	9713 (1966)	9181 (1913)	0.056	14

For abbreviations see list of abbreviations (page 6).

FIGURE 5.1. Box plots showing changes in SD-score for Total Body Fat and Fat Free Mass, 6 and 12 months after starting treatment with a PPI compared to the baseline value (at t0 → Y= 0), (*P*-values by Wilcoxon).

TABLE 5.3. Mean values and statistics of static and dynamic lung function and respiratory muscle parameters before and during PPI.

Parameter	Mean (SD)		Wilcoxon <i>P</i> =	Number
	Before PPI	During PPI		
TLC (% pred.)	109.5 (16.6)	100.3 (7.3)	0.025	13
RV (% pred.)	179.6 (73.1)	141.5 (29.9)	0.055	13
ITGV (% pred.)	128.0 (42.9)	111.0 (13.1)	0.209	13
ERV (% pred.)	71.9 (15.6)	78.6 (20.6)	0.133	13
RV/TLC (% pred.)	40.1 (10.1)	35.2 (7.2)	0.173	13
ITGV/TLC (% pred.)	55.7 (9.2)	54.0 (4.6)	0.753	13
FEV ₁ (% pred.)	83.9 (17.2)	81.8 (16.9)	0.234	14
FVC (% pred.)	86.6 (13.0)	87.6 (13.1)	0.950	14
MEF25%FVC (% pred.)	56.6 (53.7)	34.8 (16.6)	0.103	14
PI _{max} (cmH ₂ O)	-66.3 (18.9)	-81.3 (18.7)	0.002	13
PE _{max} (cmH ₂ O)	75.9 (15.8)	83.7 (18.4)	0.069	13
MIF50%FVC (% pred.)	47.3 (14.9)	47.6 (13.8)	0.925	14
sGAW (1/kPa.s)	0.864 (0.257)	0.854 (0.219)	0.807	13
PEF (% pred.)	90.6 (18.7)	87.9 (15.4)	0.177	14

For abbreviations see list of abbreviations (page 6).

FIGURE 5.2. Box plots showing changes in Maximal Inspiratory Pressure (PI_{max}) comparing each 3 month period, during treatment with a PPI, with the baseline value (at t₀ → Y=0), (*P*-values by Wilcoxon).

TABLE 5.4. Mean values and statistics of the home recordings of Peak expiratory flow (= PEF) parameters before and during PPI.

Parameter	Mean (SD)		Mean (SD)		Wilcoxon <i>P</i> =	Number
	Before PPI		During PPI			
Mean diurnal variability (l/min)	24.6	(16.7)	26.6	(15.6)	0.722	14
Maximal diurnal variability (%)	27.6	(12.3)	21.3	(4.4)	0.041	14
Minimal morning PEF (%)	72.4	(12.3)	78.4	(4.2)	0.056	14
Mean difference PEF (l/min)	5.9	(27.6)	16.5	(19.3)	0.106	14

For abbreviations see list of abbreviations (page 6).

Respiratory muscle parameters especially mean PI/PEmax improved markedly. PImax changed from 66.3 (± 18.9 cmH₂O) to 81.3 (± 18.7 ; $P=0.002$) during PPI and PEmax from 75.9 (± 15.8 cmH₂O) to 83.7 (± 18.4 ; $P=0.069$) during PPI, while MIF50%VCin did not change during PPI, from 47.3 ($\pm 14.9\%$) to 47.6 ($\pm 13.8\%$; NS) (Table 5.3 and Figure 5.2).

BRONCHIAL HYPERRESPONSIVENESS

The monthly recordings at home with the Personal Best PEF meter demonstrated variable changes. While mean diurnal variability even slightly increased from 24.6 (± 16.7 L/min) before to 26.6 (± 15.6 L/min; NS) during PPI, maximal diurnal variability decreased significantly from 27.6 ($\pm 12.3\%$ of recent best) before to 21.3 ($\pm 4.4\%$; $p<0.05$) during PPI. Minimal morning PEF increased from 72.4 ($\pm 12.3\%$) before to 78.4 ($\pm 4.2\%$) during PPI ($P=0.056$). Mean difference between best evening and best morning PEF increased from 5.9 (± 27.6 L/min) before to 16.5 (± 19.3 ; NS) under PPI (Table 5.4).

RESTING ENERGY AND DIET

REE did not change during PPI, from 111.7 % of reference values ($\pm 12.1\%$) to 110.7 ($\pm 9.1\%$; NS). Dietary parameters did not change significantly. Mean fat intake decreased from 90.5 (± 18.8 gram/day) to 81.9 (± 17.0 ; NS) during PPI and total daily energy intake decreased from 9713 (± 1966 kJ/day) to 9181 (± 1913 ; NS) during PPI (Table 5.2). The mean intake of lipase decreased from 4020 (± 970 IU/kg) to 3580 (± 880 ; NS) during PPI.

5.6. DISCUSSION

In the present study in 14 young CF patients, the effect of lansoprazole on growth, fecal fat losses, body composition, as well as the consequences of possible changes in body composition on lung function parameters and BHR was evaluated.

The decreased mean SDS for weight, height and BMI in our patients confirms the well known growth retardation of CF patients [21]. The use of a PPI resulted in a better improvement of weight SDS compared to height SDS, however, despite a visible trend, both changes were not significant when averaged for the whole treatment period. BMI, which is derived from weight and height, did improve significantly.

As this study was performed in an open designed fashion, in a small group of CF patients, the results should be interpreted cautiously. But as we in our previous open study [4], concerning the effect of a PPI on stool fat and body composition, showed exactly the same results (decrease of stool fat and increase of TBF, with unchanged FFM) a placebo effect would very unlikely be responsible for these consecutive similar findings. In the present study, performed in only partly the same population and with more than one year no use of a PPI in between, similar changes were measured and confirmed on several occasions during the treatment year. Repeated confirmation and progression of changes in the same direction during one year is more likely to be due to a treatment effect than to a placebo effect.

Our results show a significant and persistent decrease of the mean acid steatocrit during the entire study period. The finding that weight improved more than height can probably be explained by an improved fat absorption and thereby an increased TBF under treatment with a PPI. Our body composition measurements are in line with this statement. Compared to normal Dutch children our CF patients showed significant decreases of both mean TBF and FFM. Under treatment with a PPI and despite a non significant decrease in fat and energy intake, mean TBF improved significantly while mean FFM remained unchanged. The decreased steatocrit values could either be due to this slight decrease of fat intake or to improved fat absorption. Because the lipase intake decreased too, the second hypothesis seems more likely as it also explains the significant increase of body fat and BMI. These results confirm previous findings showing a favourable effect of a PPI on fecal fat losses in CF. An explanation for the diverging results concerning the improvement of FM but not FFM could be the lack of an physical exercise during the study [22].

Despite a lack of improvement of FFM we could demonstrate an improvement of the static lung volumes TLC and RV, and respiratory muscle performance, as shown by improvement of the resultant of expiratory and inspiratory forces, especially PI-max [23]. Although we could demonstrate a significant decrease in TLC and a nearly significant decrease in RV, the ratio RV/TLC appeared not to change significantly. The initial mean ratio of 40% is, however, significantly increased when compared to that of normal individuals. The lack of a significant change of ITGV is probably related to the large SD at the start. The relation between the decrease of hyperinflation (TLC and RV) and the improvement of respiratory muscle force is more difficult to interpret. Indeed a decrease of hyperinflation can lead to a more favourable starting point for diaphragm muscle contraction, improving the contractile force and therefore maximal mouth pressures [24]. Lands et al. [25] demonstrated in their comparative study, an effect of nutritional status on respiratory muscle strength in CF patients but not in asthmatic and anorexia nervosa patients, indicating that the combination of malnutrition and chronic hyperinflation give rise to inefficient functioning of the diaphragm. The

prompt decrease of TLC and RV with an increase of PI/PEmax in the present study, points to such a beneficial effect of a PPI on the diaphragm muscles. But, because of the unchanged ITGV, ERV and RV/TLC, we could not demonstrate an improved elastic recoil. In the absence of absolute changes in body FFM, the improvement of PI/PEmax during the intervention period could hypothetically be explained by metabolic adaptations in the respiratory muscles rather than by changes in the mechanical properties of the diaphragm. As ventilatory muscles training was in no way part of the intervention procedure, training effects are expected not to be related to these changes in ventilatory muscle function. Besides the PImax, the sustained maximal inspiratory pressure could have clinical significance in CF patients. Further studies are needed to assess the effects of a PPI on endurance of the inspiratory muscles.

The reduction in lung volumes under lansoprazole seems to us clinically relevant in view of the data of Kraemer et al [26], who recently showed in 60 infants with CF that differences in lung function within subgroups of mutations were mainly related to pulmonary hyperinflation. Hyperinflation seems to occur in a very early phase of the disease and may even precede inflammation and infection. This might, at least partly, be influenced by the type of mutation as shown by Kraemer.

Lung hyperinflation in CF patients have been related to surfactant deficiency [27-30]. The improved fat absorption might have a favourable effect on surfactant composition, but the relation between the use of a PPI and improvement of surfactant composition has yet to be demonstrated. Another factor contributing to hyperinflation could be the degree of inflammation, especially of the smaller airways. Bronchial inflammation is known to be present in CF, where it has been demonstrated even in very young CF patients [31].

The absence of effects on sGAW after a PPI does not exclude changes in these smaller airways. In this respect the increase in TBF under PPI intervention is remarkable. Body fat plays an important role in energy homeostasis by production of the adipocyte-derived leptin. Leptin seems to have a pleiotrophic function in the body and its administration has been shown to improve immune function [32]. Further studies are necessary in order to elucidate the relation between leptin metabolism and the observed changes in body composition in CF patients.

From the daily recordings of the PEF at home, maximal diurnal variability and minimal morning PEF are the most important indicators for the presence of BHR [19]. Both the maximal diurnal variability, being nearly 30% of recent best PEF, and the weekly minimal morning PEF, being only 72% of recent best PEF, support the assumption that BHR is increased in CF. When compared to normal children, patients with CF show an increased prevalence of BHR [33]. BHR is directly related to bronchial inflammation, but also to GER. GER is common in CF patients [34], and is often present in infants [35] and toddlers [36]. Administration of lansoprazole might consequently affect both GER and BHR. Assessment of diurnal variability of PEF has been proposed as a surrogate for measuring the degree of BHR in asthmatic patients, even when asymptomatic [37], but its usefulness has been limited because of its rather poor reproducibility [38]. However it is still the only available method to measure diurnal changes of airway obstruction in every day life [39]. We could demonstrate a significant decrease in

maximal diurnal variability and a nearly significant change in minimal morning PEF. But, on the other hand, the mean diurnal variability did not improve at all, which perhaps indicates the poor reproducibility of this method. The mean differences only slightly increased towards the end of the study, which demonstrated that evening PEF improved more than morning PEF, but this change was not significant. Although our results of PEF recordings do not allow us to conclude definitely that BHR in CF patients is decreased by antacid therapy with a PPI, our results suggest that BHR is present in CF and that GER may contribute to it. More studies are necessary to confirm this finding. In conclusion we would recommend to assess regularly, even in young CF patients, dynamic as well as static lungvolumes. Further, as our study shows that improvement of body composition and of some lung function parameters, particularly TLC and PImax occur in parallel, we would recommend to make all effort to reduce fat malabsorption in young CF patients. The relationship between body composition and lung function in children with CF should be further studied in order to elucidate the mechanisms involved.

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CHAPTER SIX

Effects of therapy with lansoprazole on intestinal permeability and inflammation in young Cystic Fibrosis patients

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6.1. ABSTRACT

Background: Defective pancreatic bicarbonate secretion with low intestinal pH or intestinal inflammation of any origin increase intestinal permeability in Cystic Fibrosis (CF).

Methods: In this open study the authors evaluated the effect of a proton-pump inhibitor (PPI) on intestinal permeability and inflammation in 14 young, pancreatic insufficient CF patients. Permeability was measured, by a three-sugar permeability test before and after 1 year lansoprazole use, and urinary nitric oxide (NO) oxidation products were assessed before and during that year, as a marker of inflammation.

Results: After 1 year of lansoprazole use, median urinary recovery percentages changed from 2.5% to 1.7% ($P=0.064$), from 24.9% to 24.5% (NS) and from 10.5% to 11.1% (NS) for lactulose, mannitol and L-rhamnose respectively. Despite the fact that the median urinary excretion ratios decreased from 0.108 to 0.083 ($P=0.03$) and from 0.246 to 0.176 ($P=0.016$) for lactulose to mannitol and lactulose to L-rhamnose, respectively, they both remained increased. Median urinary NO-products-to-creatinine ratios was 0.287 for CF patients before lansoprazole and 0.130 for healthy controls ($P=0.002$). Although there was a tendency towards a decrease in NO products-to-creatinine ratio during treatment, this was not significant at the endpoint.

Conclusions: Intestinal permeability is considerably increased in CF patients and is partly corrected after the use of a PPI for one year, which may point to a harmful effect of the acid luminal contents on the tight junctional related paracellular permeability pathway. The start and end values for the NO products-to-creatinine ratio in CF patients were not significantly different, but were considerable increased when compared to control participants ($P=0.002$).

6.2. INTRODUCTION

Cystic Fibrosis (CF) is the most common lethal hereditary disease among white persons. The CF gene is located on chromosome 7 and so far more than 900 mutations of this single gene, producing the CF transmembrane conductance regulator (CFTR), have been reported. The basic defect is a dysregulation of the chloride channels, resulting in dehydration of the luminal surface of exocrine cells. Apart from respiratory complications, gastroenterologic abnormalities give rise to significant morbidity in children with CF.

Although a severe increase in intestinal permeability (IP) is known to be present in CF patients, the clinical significance of this abnormality is unknown at present [1,2]. A recent study in rats, revealed that increases of IP favours the development of intestinal damage associated with excessive pancreatic enzyme administration. The characteristically increased IP in CF similarly could increase vulnerability to fibrosing colonopathy [3]. In a review article a relationship between increased IP, decreased intestinal pH and the pathogenesis of fibrosing colonopathy has been suggested [4]. Although a relationship has been shown to exist between IP and pancreatic insufficiency (PI), of whatever origin [5], patients with CF do show specific IP abnormalities such as increased mannitol xylose and rhamnose permeability [2,6,7]. These CF related abnormalities suggest the presence of specific pancreatic dysfunction contributing to the increased IP. One of these abnormalities could be the severe CFTR-associated decrease in ductular bicarbonate secretion [8], resulting in a low duodenal buffering capacity and therefore in a low luminal pH.

In children with CF, exogenous administration of pancreatic enzymes improved but did not reverse fat malabsorption completely. It has been shown that the use of a proton pump inhibitor (PPI) improves the effectiveness of administered pancreatic enzymes [9,10]. The purpose of these drugs is to increase duodenal pH by decreasing gastric acid production. Also, a direct action on the residual pancreatic function is possible [11], which could be more expected in young CF patients. If duodenal pH is less than 5, early dissolution of the enteric coating of administered enzyme capsules is hindered because this only occurs at a duodenal pH above 5.5 [12,13]. In addition this low pH may also inactivate pancreatic enzymes. Both processes result in suboptimal functioning of the enzymes required for the treatment of malabsorption.

Additionally, a low luminal pH could also have a direct effect on the mucosa of the intestine. As a possible explanation for the increased IP in CF, we hypothesized that the increased acidity of the intestinal contents could affect mucosal integrity. Decreased mucosal integrity could lead to both increased IP and intestinal inflammation (II), both present in CF.

Intestinal permeability is also increased in chronic inflammatory diseases such as celiac disease. In previous studies, we demonstrated an increased expression of inducible nitric oxide synthase (iNOS) in mucosal biopsies of celiac disease patients, resulting in II and high levels of urinary nitric oxide (NO) oxidation products, nitrite, and nitrate [14,15]. Prostaglandins and NO are important mediators of inflammation and there is mounting evidence for a biological relation between prostanoid and NO biosynthesis

[16]. We therefore hypothesized that NO oxidation products could be increased in CF, resulting in an increase of the NO oxidation products-to-creatinine ratios.

As part of a wider study, the aim of the present study was to investigate the effect of lansoprazole on fat malabsorption and IP and to assess the presence of II by determination of urinary NO oxidation products excretion, and to find a possible association between Intestinal permeability and inflammation. Informed consent was obtained before onset, as part of a wider study, the protocol of which was approved by the Medical Ethical Committee of the University Hospital of Maastricht.

6.3. MATERIALS AND METHODS

PATIENTS

Fourteen stable CF patients (9 males, 5-16 years of age) with PI and mild lung disease, were studied before and during 1 year of treatment with 15 mg lansoprazole (a PPI) daily. Because of this standard dose, we also computed the lansoprazole dose per kilogram of body weight. Diet and exogenous pancreatic enzymes were not changed during the intervention. For inclusion in the study, prestudy stool acid steatocrit (an index of fat losses) had to exceed 25%, despite an intake of at least 2000 IU lipase/kg bodyweight (mean 4020 IU lipase/kg). The genetic mutations have been characterized in only 8 out of 14 (57%) of our patients ($\Delta F508$ mutation in 75%). Table 6.1. summarizes the clinical data of our patients.

METHODS

Intestinal Permeability: (IP) was measured before and after 1 year of treatment with lansoprazole 15 mg daily, by 3-sugar permeability test (SPT) [17]. After an overnight fast the first urine of the day was discarded, CF patients ingested 100 ml of the test solution containing 1 g mannitol, 1 g L-rhamnose and 5 g lactulose. This solution was isotonic (osmolality 330 mmol/L). After drinking the test solution, urine was

TABLE 6.1. Patient characteristics at start

Item		Mean	Median	SD	Range
Age	(y)	9.74	8.60	3.38	4.6 – 17.2
Height	(cm)	133.9	131.5	16.6	107.0 – 162.3
	(SDS)*	-1.068	-1.112	0.590	-2.166 – 0.003
Weight	(kg)	27.7	25.6	9.9	15.5 – 52.3
	(SDS)	-1.270	-1.438	0.739	-2.393 – 0.272
Acid Steatocrit	(%)	33.3	32.7	17.7	5.4 – 65.7
FEV ₁ [†]	(% predicted)	83.9	85.0	17.2	58 – 112
FVC [‡]	(% predicted)	86.6	87.0	13.0	64 – 109

* Standard deviation score or Z score; † Forced expiratory volume in 1 second; ‡ Forced vital capacity

collected for 5 hours for determination of these 3 sugars by high-performance liquid chromatography and expressed as percentage of intake. Water intake was allowed after the first half hour in all patients, while, for practical reasons, food and sugar containing fluids intake was delayed for 3 hours. Sugar excretion ratios were determined by dividing the urinary percent recovery of the large sugar (lactulose) by that of the smaller sugars (mannitol + L-rhamnose). This simple and non-invasive test is a reproducible index of IP in children [17,18]. The use of ratios correct for potentially confounding factors such as delayed gastric emptying, delayed or rapid gut transit, or impaired renal clearance.

Nitric oxide oxidation products in urine: Nitric oxide oxidation products in urine were determined before and every 3 months during treatment. On the day before a visit, the CF patients collected urine for 24 hours, the 24 hours before and during the collection period these patients used a nitrate poor diet. Patients were free of viral illness or CF exacerbation during the collection period. A Nitrate/Nitrite colorimetric assay kit (Cayman Chemical Company, Ann Arbor, MI, US) determined nitrite and nitrate content. This test uses the Griess reagent assay, which is more accurate than the HPLC method [19]. Urinary creatinine was also determined and the NO oxidation products-to-creatinine ratio computed. The data of the CF patients were compared with urine samples of 25 healthy children (mean age 9,0 year ($\pm 4,8$), healthy children of staff members).

Fecal Acid Steatocrit: Fecal Acid Steatocrit was used to monitor the steatorrhea as previously described [20,21]. Here results are expressed as the fat layer divided by the fat plus solid layer as a percent. Fecal acid steatocrit has been shown to correlate well with chemical analyses of fecal fat ($r = 0.83$; $P < 0.001$) [22]. In healthy individuals older than 6 months, the acid steatocrit is less than 20%. We chose for this method to monitor fecal fat content repetitively and easily during the study.

Statistics: In the tables data are presented as means, standard deviations and medians, whereas in the figures the data are presented as box plots with medians and quartiles. Individual values of the 3-SPT are shown in Table 6.2. By using the non-parametric Wilcoxon signed ranks test, we avoided making an assumption of normality, which is untestable with only 14 subjects. Differences with respect to baseline were considered significant when the two-sided P -value was 0.05 or less. The statistical package SPSS (version 9.0 for Windows, 1999, SPSS Inc., Microsoft, Chicago, IL) was used for data management and analysis.

6.4. RESULTS

INTESTINAL PERMEABILITY

Intestinal Permeability was severely increased in all but one subject before adjuvant therapy with lansoprazole. After 1 year of treatment with lansoprazole, the

TABLE 6.2. Case Summaries of fat losses (acid steatocrit) and 3 sugar permeability test, before and after lansoprazole for one year

Case	Steatocrit (%)		L-rec.*(%)		M-rec.†(%)		Rh-rec.‡(%)		L/M¶ (ratio)		L/Rh§ (ratio)	
	Start	End	Start	End	Start	End	Start	End	Start	End	Start	End
1	42.2	39.0	2.5	1.5	18.1	21.7	7.8	8.5	0.14	0.07	0.32	0.18
2	31.2	23.0	7.2	1.7	7.4	8.8	9.7	7.3	0.98	0.20	0.75	0.24
3	38.5	26.7	2.2	.5	25.1	31.9	10.6	4.0	0.09	0.02	0.21	0.12
4	26.3	11.6	7.4	2.2	24.8	24.8	12.8	12.4	0.30	0.09	0.58	0.18
5	65.7	38.6	3.2	1.9	27.6	35.2	11.8	16.8	0.12	0.05	0.27	0.11
6	29.0	16.4	4.6	1.9	21.4	19.8	10.5	11.0	0.21	0.10	0.43	0.17
7	5.4	24.6	2.4	1.7	17.0	17.6	8.9	10.8	0.14	0.10	0.27	0.16
8	14.1	15.8	.8	5.8	49.7	40.6	10.5	18.7	0.02	0.14	0.08	0.31
9	7.8	8.9	4.5	2.1	32.2	21.9	18.5	12.5	0.14	0.10	0.25	0.17
10	38.3	37.2	2.6	1.3	32.9	25.3	10.3	7.4	0.08	0.05	0.25	0.18
11	34.2	41.1	2.7	1.6	36.5	35.3	16.1	15.1	0.07	0.05	0.17	0.11
12	45.3	18.9	2.1	4.1	20.9	30.1	10.2	16.7	0.10	0.13	0.20	0.24
13	25.6	39.2	.9	1.7	12.4	21.7	5.2	9.1	0.07	0.08	0.18	0.18
14	62.3	35.7	1.9	1.0	26.3	24.2	11.4	11.1	0.07	0.04	0.17	0.09
Mean	33.3	26.9	3.2	2.1	25.2	25.6	11.0	11.5	0.181	0.087	0.294	0.175
Median	32.7	25.6	2.5	1.7	24.9	24.5	10.5	11.1	0.108	0.083	0.246	0.176
SD	17.8	11.4	2.0	1.3	10.7	8.3	3.3	4.2	0.240	0.048	0.180	0.059
Normal values £			0.11 – 0.67		8.1 – 20.2		4.1 – 10.9		0.011 – 0.036		0.023 – 0.074	

* Lactulose recovery; † Mannitol recovery; ‡ L-rhamnose recovery; ¶ Lactulose-to-mannitol ratio; § Lactulose-to-L-rhamnose ratio; £ reference range according to Miki et al (17).

TABLE 6.3. Three-sugar test: change in recovery and excretion ratios after treatment with lansoprazole for 1 year

	Lactulose recovery	Mannitol recovery	Rhamnose recovery	L/M ratio (% recovery)	L/Rh ratio (% recovery)
Mean change	- 1.138	+ 0.467	+ 0.504	- 1.015	- 0.913
Mean % decrease	35.4	- 1.9	- 4.5	52.0	40.5
Median change	- 1.174	+ 0.310	+ 0.132	- 0.707	- 0.670
SD	2.64	6.46	4.34	1.30	0.92
Wilcoxon (P =)	0.064	0.826	0.730	0.030	0.016

L, lactulose; M, mannitol; Rh, L-rhamnose

FIGURE 6.1. Box plots of urinary recovery of lactulose, mannitol and L-rhamnose before and after 1 year treatment with a proton-pump inhibitor, expressed in % of administered dose (dotted lines are reference values). P values were determined by Wilcoxon rank sum test.

FIGURE 6.2. Box plots of urinary recovery ratios of large (lactulose) versus small sugars (mannitol and rhamnose) before and after 1-year treatment with a proton-pump inhibitor. Dotted lines are maximum of reference values. P values were determined by Wilcoxon rank sum test.

TABLE 6.4. Significance of bivariate correlations of lansoprazole/body weight with lactulose recovery, NO oxidation products/creatinine, L/M and L/Rh ratios

Lansoprazole/ body weight	NO oxidation products/ creatinine	L/M ratio	L/Rh ratio	L recovery
start	0.608	0.805	0.454	0.655
12 months	0.244	0.961	0.782	0.308

L, lactulose; M, mannitol; Rh, L-rhamnose

TABLE 6.5. Mean and Median values of the nitric oxide oxidation products-to-creatinine ratio (mmol/mmol) before and during 1 year use of lansoprazole

	Before	3 months	6 months	9 months	12 months	Controls
Mean	0.330	0.326	0.261	0.216	0.270	0.166
Median	0.287	0.232	0.235	0.233	0.252	0.130
SD	0.186	0.228	0.150	0.102	0.101	0.114
P value *		0.470	0.594	0.019	0.140	0.002

* Wilcoxon rank sum test compared to value at start

mean urinary recovery percentages changed from 3.22% to 2.08%, from 25.16% to 25.62% and from 11.02% to 11.53% for lactulose, mannitol, and L-rhamnose, respectively. Lactulose recovery decreased by 35%, from five times to three times the upper reference limit, but the decrease just failed to reach statistical significance ($P = 0.064$). The urinary excretion ratios decreased from 0.181 to 0.087 ($P = 0.03$) and from 0.294 to 0.175 ($P = 0.016$) for lactulose to mannitol and lactulose to L-rhamnose, respectively. Although both ratios decreased by half, they still were more than double the upper reference limit (Tables 6.2. + 6.3., figure 6.1. + 6.2.).

FECAL ACID STEATOCRIT

Fecal Acid Steatocrit decreased significantly from a mean of 33.3% ($\pm 17.8\%$) at the start to 23.7% ($\pm 11.0\%$) at 3 months ($P < 0.01$) and remained at that level thereafter, the mean value during treatment was 24.0% (9.8%; $P < 0.05$). Even when converted for body weight, no correlations were found between lactulose recovery, lactulose to mannitol and lactulose to L-rhamnose ratios and the mean fecal acid steatocrit, as an index for persisting fat losses, however (Table 6.4.).

URINARY NITRIC OXIDE OXIDATION PRODUCTS

Urinary nitric oxide oxidation products expressed as median (SD) NO oxidation products-to-creatinine ratio in CF patients was, at the start of PPI treatment, 0.287 (± 0.186) whereas it was 0.130 (± 0.114) for control participants ($P = 0.002$). There was a trend in decrease of the mean NO oxidation products-to-creatinine ratio during the treatment period, with 0.233 (± 0.102) as the lowest value at 9 months ($P = 0.019$), which was still significantly increased compared to the healthy controls

($P=0.035$). The value at the end, however, was increased again and was not different from start values (Table 6.5.).

6.5. DISCUSSION

In this open prospective study, we demonstrated that IP was dramatically increased in CF and more pronounced for the lactulose recovery. After the use of lansoprazole for 1 year, the 3-SPT only showed an improvement in the permeability of the large sugar, lactulose. Therefore both ratios lactulose-to-mannitol and lactulose-to-L-rhamnose ratios decreased significantly. During that year, the NO oxidation products-to-creatinine ratio tended to decrease, but was in the end not significantly different from start values. Furthermore, after the use of a PPI for 1 year in these same patients, the fecal acid steatocrit decreased and total body fat improved significantly ($P < 0.05$), whereas free fat mass remained unchanged. SD-scores of height and body weight showed a slight, but non significant, increase [23]. Because this study was performed with an open design in a small group of CF patients, the results should be interpreted cautiously. In the present study, we only tried to determine the change in increased IP before and after 1 year of PPI treatment. For ethical reasons, we did not check IP in the healthy control group, before and after 1 year of lansoprazole use. Although we do not expect that lansoprazole could improve IP by itself, we are not certain about this. Nevertheless we consider the observed improvement in IP likely to be linked to a decrease in luminal acidity.

Although it is well known that IP is increased dramatically in CF [1-7], the cause of this phenomenon is poorly understood. Van Elburg et al. [24] have shown the increased IP in CF to be linked to pancreatic insufficiency and not to the dose of pancreatic enzyme supplementation. However, IP in CF has been related to genotype, and therefore more basic defects[7]. We hypothesized that increased acidity, resulting from the CFTR-associated decreased bicarbonate secretion, might affect the small bowel mucosal integrity. Although our study was not placebo controlled, we could demonstrate after intervention with lansoprazole for 1 year a decrease in IP for the larger sugar lactulose only, whereas the recoveries of the smaller sugars remained unchanged. This contrast in effects leads to a significant decrease of lactulose-to-mannitol ($P = 0.03$) and lactulose-to-L-rhamnose ($P = 0.016$) excretion ratios. According to the "pore theory", permeation takes place through paracellular pathways (large pores) and transcellular pathways (small pores)[25]. In the 3-SPT, the sugars permeate via different routes, L-rhamnose and mannitol predominantly transcellularly and lactulose as a large sugar only permeates paracellularly. Because in our patients only lactulose recovery improved, our results indicate in duplo that adjuvant therapy with PPI only affects the paracellular or "tight junctional" pathway. As to the cause of the increased IP in CF, it could be either primary or secondary. Primary tight junctional abnormalities at the level of the pancreas, ileum and trachea in CF patients have been reported previously [26]. A possible direct relation between

the basic cellular CF defect and the increased intestinal permeability and inflammation has recently been suggested [27,28].

Low intestinal pH resulting from defective pancreatic bicarbonate secretion or II of any origin belongs to possible secondary causes of increased IP in CF. First, low intestinal pH either could damage tight junctions directly or be responsible for late opening of the enzyme coating in the small bowel, resulting in elevated luminal proteolytic activity, leading to tight junctional damage and increased IP. In both cases one would expect IP to decrease after use of a PPI, as shown by our results. The increased IP did not normalize probably because PPIs only slightly increase distal duodenal pH, but do not normalize it [9,10,29]. Second, II of various origin such as gastroenteritis, inflammatory bowel diseases, and celiac disease, are known to be associated with increased IP and could be responsible for or could aggravate the increased IP in CF [30]. Although the bowel mucosal architecture is usually considered to be normal in patients with CF [31], mucosal inflammation has been shown to be present [27,28]. Like other researchers we showed that the urinary NO oxidation products-to-creatinine ratio was increased in acute celiac disease [14,32]. Therefore we evaluated this marker of cytokine-induced increased NO synthesis by measuring urinary NO oxidation products in our CF patients. The NO oxidation products-to-creatinine ratio found in the CF patients proved to be significantly increased ($P=0.002$), indicating that inflammation could be present. After use of a PPI for 1 year, this ratio tended to decrease, but this decrease at the end was not significantly different from start values. We checked for possible causes, such as infections, changes in compliance to therapy, increase in weight and diarrhea, that could not explain the observed variations in the NO oxidation products-to-creatinine ratios. Although regression to the mean could be responsible for the apparent variations in NO oxidation products-to-creatinine ratio, these variations could also reflect variations in inflammation at both intestinal and pulmonary levels [33]. Third, although IP improved, it did not normalize, which could reflect persisting II and elevated NO oxidation products-to-creatinine ratios. Fourth the possibility remains that 15 mg lansoprazole daily for 1 year in children with CF is not sufficient to normalize intestinal permeability and inflammation, because it also did not normalize fat malabsorption [23]. Therefore our results do not allow any firm conclusions about the possible relationship between intestinal permeability and inflammation. Further studies are necessary to elucidate this point.

Finally, during the treatment year and in the years thereafter, patients did not report adverse events, such as diarrhoea, constipation, or colitis. No vitamin B₁₂ deficiency could be demonstrated, even after 3 years of use of PPI [34].

In conclusion, the use of a PPI as adjuvant therapy in children with CF resulted in improvement of fat malabsorption and IP, linked to a significant decrease of the paracellular permeability pathway only. Because PPI therapy was the only intervention, we consider the increased acidity of intestinal contents to be a likely cause of increased IP. The introduction of a PPI improved but did not normalize IP. We also could demonstrate a significant increase of urinary NO oxidation products-to-creatinine ratios, pointing to the presence of inflammation in our patients, possibly of intestinal origin. But because of lack of difference at the end point, we are

not able to draw any conclusion about therapy with lansoprazole and a possible link between increased intestinal permeability and inflammation.

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CHAPTER SEVEN

Fatty acid status of plasma and red blood cells phospholipids in Cystic Fibrosis patients treated with and without lansoprazole

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7.1. ABSTRACT

Background: In a former study on the additional effect of a proton pump inhibitor (PPI) in Cystic Fibrosis (CF) patients, we were able to demonstrate an improvement in fat absorption.

Objective: The present study was performed to assess the (essential) fatty acid ((E)FA) composition of plasma and red blood cells (RBC) phospholipids (PL) in patients with CF. Secondly the effect of longterm treatment with a PPI on the FA status was examined.

Design: This descriptive study compares in a cross-sectional design, 11 CF patients who used lansoprazole (15 mg/day) for 3 years (PPI+ group), 9 CF patients on the standard regime (PPI- group) and 10 healthy controls. There was no baseline investigation prior to PPI treatment. Student-t test was used for comparisons of FA results between the 3 groups.

Results: Mean plasma docosahexaenoic acid (DHA) concentrations in the PPI- group (14 mg/L) and in the PPI+ group (21 mg/L), were significantly decreased ($P=0.001$ resp. $P<0.05$) compared to controls (33 mg/L). The difference between the two CF groups was significant also ($P<0.05$). Arachidonic acid (AA)/DHA and mead acid/AA ratios were significantly increased in the PPI- group ($P<0.01$). In RBC only the FA status linolenic acid was decreased and eicosatrienoic acid increased, while DHA and AA levels were normal.

Conclusion: Our CF patients showed biochemical signs of EFA deficiency mainly in plasma PL. Although the EFA status of patients receiving a PPI, showed a trend towards normalization, the only significant difference between the two CF groups consisted in a higher plasma DHA level in the PPI+ group. Our results suggest some role for fat malabsorption in the different EFA status of CF patients.

7.2. INTRODUCTION

Cystic Fibrosis (CF) is an autosomal recessive inherited disorder, characterised by a defect in the chloride transport regulating channel at the luminal side of exocrine cells, the cystic fibrosis transmembranous conductance regulator protein (CFTR). One of the most serious consequences in CF patients of this defect is pancreatic insufficiency, which induces malabsorption, especially for fat. This pancreatic insufficiency is only partially corrected by exogenous pancreatic enzymes [1]. The malabsorption in CF enhances the poor nutritional status seen in these patients [2]. Fat nutritional status can be biochemically assessed, for instance by analysing fatty acid (FA) profiles [3]. Several studies have shown the presence of an essential fatty acid (EFA) deficiency or shortage in CF patients [3-7]. However, controversy exists about the origin of this EFA shortage [8-13]. A few reports have been published on the lower EFA status of red blood cell phospholipids (RBCPL) [14-16]. A complete FA profile of RBCPL in CF has not yet been documented in complete detail. Recent studies demonstrated that, in a CFTR knockout mice model, lipid membranes of CF affected organs, such as ileum, pancreas, and lung, show a profound decrease of docosahexaenoic acid (DHA, 22:6n-3) and an increase of arachidonic acid (AA, 20:4n-6). These abnormalities could be reversed by oral administration of DHA [17,18].

Although even in patients with a normal EFA status lung disease will develop, it has been postulated that an EFA shortage could be a predisposing factor for lung disease in CF patients [19,20]. An improvement of the nutritional status with optimized EFA status could lead to a decrease of inflammation and an improvement of lung function. In two earlier studies we have shown that lansoprazole, a proton pump inhibitor (PPI), used as adjuvant therapy, significantly improved both steatorrhea and the nutritional status of CF patients while at the same time lung hyperinflation decreased in those patients with a mean age of 10 years [21,20]. The effect of a PPI on the EFA status of CF patients has not been reported so far. The present study was performed primarily to assess the detailed FA composition of plasma and RBCPL in CF patients. In addition, the EFA status of CF patients using a PPI for 3 years was compared to that of CF patients without a PPI and that of control subjects.

7.3. SUBJECTS

Twenty CF patients from our out-patient clinic were studied. All patients had the diagnosis confirmed by sweat chloride testing (Chloride more than 60 mmol/l) and a persistent steatorrhea which required regular pancreatic enzyme replacement therapy. Eleven of these patients (PPI+ group, 6 males) were recruited earlier for a one-year study evaluating the effects of a PPI on fat malabsorption and body composition. The selection criteria for that study consisted of persisting steatorrhea, a forced expiratory volume in 1 second (FEV₁) of more than 70% of predicted, and stable CF disease [22]. The observed improvement was the reason for continuing the PPI therapy. They received lansoprazole treatment in a dose of 15 mg/day for at least 3

years as adjuvant treatment to the standard regime consisting of exogenous pancreas enzyme replacement therapy, an energy and fat enriched diet, vitamins A, D, E and K supplements, and nebulization of mucolytics and chest physiotherapy. The mean age of this group at the start of lansoprazole therapy was 8.3 years (range: 5.3-14.0) and at the time of FA analysis 11.7 years (8.5-17.4). In this group DNA-analysis had taken place in four patients. Three patients were homozygous for the $\Delta F508$ mutation. One subject had a $\Delta F508/N1303K$ mutation. The remaining 9 patients (PPI- group, 4 males) received standard treatment as described above. In this group the mean age at the time of FA analysis was 8.2 years (1.8-12.3). DNA-analysis were performed in seven out of these nine patients. Four patients were homozygous for the $\Delta F508$ mutation. The other 3 patients were heterozygous for $\Delta F508$.

All patients were screened for fat malabsorption at the start of the study by measuring faecal acid steatocrit values [23]. Our upper range of normal for acid steatocrit is 20%. In the PPI+ group the mean steatocrit value before PPI treatment was 41.7% (range: 22.4-65.7%) after 3 years, at the time of sampling for FA analysis, the acid steatocrit was significantly decreased to a mean of 28.6% ($\pm 12.2\%$). The PPI- group had a mean steatocrit value of 32.6% (range: 7.9-67.2%). The difference in mean steatocrit values between the two CF groups was not significant. In the PPI+ group 2 patients suffered from a meconium ileus at birth whereas this was the case for 3 patients in the PPI- group.

The two CF groups described above were compared with a healthy control group consisting of ten preoperative surgical patients (6 males), not suffering from any inflammatory, infectious, digestive or nutritional disease. The mean age of the control group at FA analysis was 9.2 years (3.3-16.2).

Informed consent was obtained from all the parents and from children older than 11 years, before onset of the study. The study protocol was approved by the medical ethical committee of the university hospital of Maastricht.

7.4. METHODS

The fatty acid profiles were determined in plasma and RBCPL as previously described [24]. Briefly, lipid extracts were prepared from both plasma and RBC samples, and the phospholipid fraction separated using aminopropyl-BondElut cartridges [9]. The phospholipid FA were hydrolyzed and transmethyalted with boron trifluoride in methanol. The composition of the obtained FA methyl esters was determined by capillary gas chromatography using a polar column (50 m x 0.25 mm CPSil 88; Chrompack, Middelburg, the Netherlands), with helium as carrier gas, pressure 130 kPa. Both the injection and the detector temperatures were 300° C. The starting temperature of the column was 160° C. After 10 min, the temperature increased to 190° C with a rate of 3.2° C/min. Subsequently, after 15 min, the temperature was further increased to 230° C with a rate of 5° C/min. The amount of each fatty acid was quantified by the amount recovered of the internal standard (dinonadecanoyl [19:0]-phosphatidylcholine) added to each sample.

7.5. STATISTICAL ANALYSIS

Statistical analysis of the relative data (expressed as % of total amount of FA) and the absolute data were not different. Because of the debate whether or not absolute numbers expresses better real life necessity of EFA, we presented them both. In the tables FA composition of plasma and RBCPL are presented as absolute values in mg/L (mean SD). Relative concentrations are shown in figure 2, while in figure 3 box plots with medians and quartiles of the absolute concentrations of DHA are presented. Because the FA data distribution did not depart from normality, the separate variances Student t-test was used for comparisons between groups ($\alpha=0.05$, 2-sided). For statistical analysis the SPSS package for windows (version 10.0) was used.

7.6. RESULTS

PLASMA PHOSPHOLIPIDS

FA concentrations of plasma phospholipids (PLPL) of the PPI+ group, the PPI- group and the control group are shown in table 7.1. and summarised in figure 7.1. In the PPI- group the total FA concentration of PLPL was slightly but significantly ($P<0.05$) lower compared to the concentrations found in the PPI+ and control groups. As table 1 shows, most items of the FA phospholipids (for PPI- group 14/24, for PPI+ group 6/24) were significantly different from our control results.

PPI- group: Besides the total FA, mean concentrations of palmitic (16:0), linoleic (18:2n-6), docosapentaenoic (22:5n-3) and docosahexaenoic acid (22:6n-3) in PLPL of the PPI- group were lower compared to control subjects. On the other hand, mean concentrations of eicosatrienoic or mead acid (20:3n-9) and γ -linolenic acid (18:3n-6) were higher in the PPI- group compared to the control group. We also calculated the sum and ratios of different FA. These results are also shown in table 7.1. The sums of both n-3 and n-6 FA in PLPL were significantly lower, while the long chain polyunsaturated (lcp) n-9 FA was significantly higher in the PPI- group compared to control subjects. Furthermore, significantly lower amounts of saturated FA (SFA) and polyunsaturated FA (PUFA) were observed of PLPL of the PPI- group. The ratio of 20:3n-9/20:4n-6, which is a generally accepted marker for EFA shortage [25], showed to be higher in PLPL of the PPI- group compared to control subjects but the critical value for EFA shortage (>0.2) was not reached (figure 7.2.). The ratio 20:4n-6/22:6n-3 was also significantly higher.

PPI+ group: In contrast to the results of the PPI- group, the results of the PPI+ group were less different in comparison to the control group. No significant differences were found for mean concentrations of 16:0, 18:2n-6, 20:3n-9, total n-6, total n-9lcp, SFA, PUFA and total FA in PLPL. When compared to concentrations of our control group, the mean concentrations of 18:3n-6, 22:5n-3, 22:6n-3, total n-3, and the ratios 20:3n-9/20:4n-6 and 20:4n-6/22:6n-3 in PLPL were slightly but significantly different, with values between those of the control and the PPI- group.

FIGURE 7.1. Schematic view of fatty acid synthesis, showing different findings in plasma phospholipid fatty acids (FA). Black arrows indicate significant changes of FA concentrations in CF patients treated without lansoprazole (PPI–) compared to control subjects. White arrows indicate significant changes of FA concentrations in CF patients treated with lansoprazole (PPI+) compared to control subjects. The significance of differences with the control group are shown by small arrows ($P<0.05$) and large arrows ($P<0.01$). Differences between the two CF groups is shown by ‡ ($P<0.05$). All data are computed by student t-test.

No differences between the three investigated groups were found in mean concentrations of 16:1n-7, 18:0, 18:1n-9, 18:3n-3, 20:3n-6, 20:4n-6, 20:5n-3, 22:4n-6, 22:5n-6, and MUFA.

When comparing the PPI+ group to the PPI– group only the DHA and AA/DHA proved to be significantly different ($P<0.05$), while the mead acid/AA ratio, as a marker of EFA deficiency, was not significantly different (table 7.1. and figure 7.2.+7.3.).

RED BLOOD CELL PHOSPHOLIPIDS

Results of FA analysis of RBCPL are shown in table 7.2. In RBC, FA compositions were less abnormal when compared to plasma findings in both investigated CF groups. Most of the results of the PPI+ group were in between those of the PPI– group

FIGURE 7.2. Mean (+SD) relative plasma: levels of arachidonic acid (AA), docosahexaenoic acid (DHA) and mead acid / arachidonic acid ratio, in CF patients without (PPI-) and with (PPI+) proton pump inhibitor treatment compared to healthy controls. Statistics: both CF groups compared to controls by student t-test. (NS: not significant)

FIGURE 7.3. Box plots in medians and quartiles of absolute docosahexaenoic acid (DHA) concentrations in plasma, in CF patients without PPI (PPI-), with PPI (PPI+) and in healthy controls. Statistics of parametric (Student t-test) = non parametric (Wilcoxon) test.

TABLE 7.1. Mean absolute concentrations ($\pm$ SD) of Fatty Acids in plasma phospholipids (mg/L)

Fatty Acids	Control subjects (n=10)	CF patients without lansoprazole (PPI-) (n=9)	CF patients with lansoprazole (PPI+) (n=11)
Total FA	1080.1 (187.4)	880.4 (154.6) ³	971.8 (283.7)
16:0	283.6 (70.1)	213.6 (41.9) ³	258.0 (98.3)
18:0	146.6 (21.4)	134.6 (19.7)	132.1 (24.2)
Total SFA ¹	482.4 (90.8)	383.7 (64.9) ³	427.8 (123.9)
16:1n-7	4.3 (1.5)	5.5 (2.6)	6.5 (4.1)
18:1n-9	94.4 (19.2)	95.9 (25.2)	99.6 (39.8)
20:3n-9	0.9 (0.6)	2.6 (1.9) ³	1.9 (2.0)
Total n-9 lcp [†]	1.6 (0.7)	3.7 (2.3) ³	2.8 (2.5)
18:2n-6	234.7 (41.5)	183.4 (32.9) ²	202.1 (71.9)
18:3n-6	0.6 (0.4)	1.5 (1.1) ³	1.2 (0.6) ²
20:3n-6	34.3 (8.1)	37.0 (11.6)	41.5 (19.4)
20:4n-6	95.8 (23.4)	76.6 (20.0)	81.1 (21.6)
22:4n-6	4.1 (0.7)	4.0 (1.4)	3.4 (0.7)
22:5n-6	2.7 (1.0)	2.6 (1.3)	2.6 (1.3)
Total n-6	377.7 (63.8)	310.0 (53.2) ³	337.9 (105.5)
18:3n-3	1.7 (0.6)	1.7 (1.0)	1.9 (1.1)
20:5n-3	5.7 (5.4)	4.1 (2.2)	4.3 (1.9)
22:5n-3	9.2 (2.1)	5.9 (2.0) ²	6.6 (2.7) ³
22:6n-3	32.9 (13.2)	13.8 (3.5) ^{2,*}	21.4 (9.7) ^{3,*}
Total n-3	50.7 (17.8)	26.5 (8.0) ²	35.1 (12.8) ³
Total MUFA ¹	138.6 (26.9)	133.3 (32.0)	144.9 (53.3)
Total PUFA ¹	430.0 (75.2)	340.2 (60.6) ³	375.9 (114.9)
20:3n-9 / 20:4n-6	0.009 (0.005)	0.036 (0.025) ²	0.026 (0.026) ³
20:4n-6 / 22:6n-3	3.1 (0.8)	5.7 (1.6) ^{1,*}	4.2 (1.1) ^{3,*}
22:5n-6 / 22:4n-6	0.65 (0.15)	0.64 (0.22)	0.74 (0.25)
22:6n-3 / 22:5n-6	14.3 (8.7)	7.9 (8.2)	8.8 (4.1)

[†] lcp: long chain polyunsaturated FA; SFA: saturated FA; MUFA: monounsaturated FA; PUFA: polyunsaturated FA.

¹⁻³ Differences versus control subjects (student-t: 1 $P < 0.001$, 2 $P < 0.01$, 3 $P < 0.05$).

* Differences between the two CF groups (student-t: $P < 0.05$).

TABLE 7.2. Mean absolute concentrations ($\pm$ SD) of Fatty Acids in red blood cell phospholipids (mg/L)

Fatty Acid	Control subjects (n=10)	CF patients without lansoprazole (PPI-) (n=9)	CF patients with lansoprazole (PPI+) (n=11)
Total FA	1107.8 (160.5)	1098.5 (89.6)	1069.8 (152.9)
16:0	248.5 (34.2)	249.3 (21.9)	256.4 (38.2)
18:0	97.4 (14.8)	94.2 (8.7)	83.8 (15.3)
Total SFAs [†]	443.2 (62.7)	432.0 (34.5)	424.6 (60.0)
16:1n-7	3.1 (1.1)	4.6 (1.6) ³	4.8 (2.0) ³
18:1n-9	139.6 (27.0)	143.6 (23.3)	140.1 (26.0)
20:3n-9	0.3 (0.3)	1.2 (0.7) ²	0.7 (0.5) ³
Total n-9 lcp [†]	0.8 (0.3)	1.8 (0.8) ²	1.2 (0.7)
18:2n-6	129.9 (23.4)	110.0 (14.2) ³	113.8 (26.4)
18:3n-6	0.5 (0.4)	0.7 (0.3)	0.6 (0.3)
20:3n-6	16.3 (4.1)	20.9 (7.3)	19.6 (5.3)
20:4n-6	116.4 (15.2)	120.7 (9.1)	112.1 (19.1)
22:4n-6	30.6 (5.5)	32.6 (6.3)	27.8 (7.2)
22:5n-6	3.5 (1.0)	4.9 (1.6)	3.8 (1.0)
Total n-6	307.2 (42.1)	301.8 (25.1)	289.8 (46.7)
18:3n-3	1.4 (0.6)	1.4 (0.8)	1.3 (0.5)
20:5n-3	3.3 (1.5)	3.8 (0.9)	3.8 (1.5)
22:5n-3	17.6 (3.0)	17.4 (2.4)	16.5 (5.1)
22:6n-3	27.7 (8.1)	22.2 (3.9)	23.6 (6.7)
Total n-3	50.4 (11.1)	45.5 (5.5)	45.6 (9.1)
Total MUFAs [†]	206.7 (38.2)	224.2 (43.6)	215.4 (42.5)
Total PUFAs [†]	358.3 (48.7)	349.1 (26.6)	336.6 (49.8)
20:3n-9 / 20:4n-6	0.002 (0.002)	0.010 (0.006) ²	0.006 (0.005) ³
20:4n-6 / 22:6n-3	4.5 (1.1)	5.6 (1.1)	5.1 (1.4)
22:5n-6 / 22:4n-6	0.12 (0.03)	0.16 (0.05)	0.14 (0.05)
22:6n-3 / 22:5n-6	8.9 (4.6)	5.1 (2.2)	6.5 (2.6)

[†] lcp: long chain polyunsaturated FA; SFA: saturated FA; MUFA: monounsaturated FA; PUFA: polyunsaturated FA.

¹⁻³ Differences versus control subjects: ¹ $P < 0.001$, ² $P < 0.01$, ³ $P < 0.05$.

and the control group. The total RBCPL FA concentration did not differ between the 3 groups.

PPI- group: In RBCPL of the PPI- group a lower concentration of 18:2n-6 was observed. On the other hand, these CF patients showed higher amounts of 16:1n-7, 20:3n-9 and n-9lcp FA in RBCPL when compared to control subjects. Further, a higher ratio of 20:3n-9/20:4n-6 was found in RBCPL in the PPI- group.

PPI+ group: In the PPI+ group no difference was seen in the concentrations of 18:2n-6 and n-9lcp compared to the control group. Only the concentrations of 16:1n-7, 20:3n-9 and the ratio of 20:3n-9/20:4n-6 were slightly higher in the PPI+ group when compared to the control group.

When comparing the PPI+ group to the PPI- group no significant differences were observed, neither for separate FA, nor for the various ratios.

7.7. DISCUSSION

In agreement with other studies [3-7], our data show a different FA profile in both plasma and RBCPL in CF patients who received the standard treatment. Although the design of the study was not adequate to compare the results of addition of a PPI for 3 years to the standard treatment, the results showed a trend towards normalisation in the PPI+ group for most FA, especially in plasma.

The evidence for an EFA shortage in our PPI- group consists of a higher concentration of 20:3n-9 (mead acid) and the 20:3n-9/20:4n-6 ratio, and lower 18:2n-6, 22:5n-3, 22:6n-3 (DHA), total n-6 FA and total n-3 FA concentrations in PLPL. Although most of our results are in good agreement with data from the literature, some differences were seen. In contrast with our results, a decrease in 20:4n-6 (AA) was demonstrated by Benabdeslam et al [3] in plasma and by Biggemann et al [15] in RBC. Like Biggemann and Freedman [18, 19], we could demonstrate a very significantly lowered DHA, especially in the PPI- group ($P=0.001$). Moreover, despite a significantly lower plasma DHA in the PPI+ group, compared to the control group, DHA was significantly higher than in the PPI- group ($P<0.05$; see table 7.1. and 7.2., figure 7.1.-7.3.). It is also important to notice that 20:3n-9, a marker of EFA deficiency, was significantly higher in the PPI- group, both in plasma as well as in RBC. Hubbard et al. [16] also found an increase of 20:3n-9 in RBCPL in pancreatic insufficient but not in pancreatic sufficient CF patients. The FA patterns in plasma and RBC are reasonably comparable to the results of healthy controls, and are suggestive of an EFA shortage. In our PPI+ group, the EFA status remained slightly disturbed, although evidently less than that of the PPI- group. The wide range of concentrations and the small number of patients could be responsible for the fact that the differences in EFA, found between both CF groups, only rarely reached statistical significance.

The less optimal EFA status, observed especially in the PPI- group, is probably due to various causes. The most frequently postulated mechanism is that EFA shortage is secondary to malabsorption of dietary fat due to pancreatic insufficiency [8]. Increased FA oxidation due to elevated energy requirements in CF patients is a plau-

sible alternative [12]. The EFA shortage could also be explained by a defect in FA metabolism [9], peroxysomal dysfunction [13] or increased lipid turnover in cell membranes [10]. It also has been shown to increase eicosanoid production [11]. Recently it has been suggested that the CFTR itself might play a role in membrane phospholipids differences in ileum, pancreas and lung, organs typically affected in CF. In these studies oral administration of DHA in CF knockout mice reversed these lipid differences [17, 18]. Differences in EFA status between the PPI- and PPI+ group could be explained in various ways. Lloyd-Still et al [20] showed EFA deficiency to be more pronounced in CF patients who had a meconium ileus at birth. We looked for this confounder and found a similar incidence of meconium ileus in both investigated CF groups, which excludes this potential explanation. Alternatively, it could be explained by a difference in CF pheno- or genotypes between the PPI- and PPI+ group. Unfortunately the genotype was only determined in about half of our studied CF patients.

Although we should be cautious in drawing definitive conclusions from this unrandomised, descriptive study, our findings show a less severe EFA shortage in the PPI+ group, as shown by the decrease of mead acid and the 20:3n-9/20:4n-6 ratio. This suggests a potential beneficial effect of lansoprazole on the EFA status in pancreatic insufficient CF patients. As lansoprazole significantly improved both steatorrhea and body composition in CF patients [21, 22], the present study might support the hypothesis that the marginal EFA status in CF is, at least partly, related to fat malabsorption. This hypothesis is further supported by the finding that the EFA status of CF patients without fat malabsorption is usually better than of patients with fat malabsorption [8,26,27]. Despite treatment with pancreatic enzymes, residual fat malabsorption in CF could be explained by a low duodenal pH associated with CF resulting in either a poor release of enteric coated pancreatic enzymes, a decreased lipolytic activity, or bile acid precipitation [28-30]. Lastly Kalivianakis et al [31] demonstrated a diminished absorption of long-chain FA through the enterocyte in CF independently of lipolytic enzyme activity. By increasing the luminal pH, a PPI could improve the solubilization and absorption of long-chain FA.

In a mouse model of CF, administration of 18:3n-3 resulted in a further decrease in DHA [18]. This suggests that there might be a block in the biosynthesis of 22:5n-3 and 22:6n-3.

In general, when DHA is low, the conversion of 22:4n-6 into 22:5n-6 becomes increased [32]. In our CF patients neither the amounts of 22:5n-6 nor the ratio between 22:5n-6 and its precursor 22:4n-6 were increased as compared to controls. This suggests a reduced activity of the metabolic processes involved in the endogenous synthesis of both DHA as well of its 'shortage marker' 22:5n-6 from their respective precursors 22:5n-3 and 22:4n-6. This synthetic pathway includes elongation of 22:5n-3 and 22:4n-6, $\Delta 6$ -desaturation of the resulting FA (both occurring in the microsomes) and 1 cycle of peroxisomal β -oxidation [33]. A reduced $\Delta 6$ -desaturation is rather unlikely, since in CF patients a lower linoleic acid status is associated with higher amounts of 18:3n-6, the $\Delta 6$ -desaturation product of linoleic acid. It can not be excluded, however, that the $\Delta 6$ -desaturase activities are mediated by different enzymes, of which one can be defective in CF. A reduced peroxisomal FA oxidation is

another option to explain the lower DHA status in CF patients without a concomitant reactive increase in 22:5n-6. This hypothesis seems the more attractive since peroxisomal dysfunction has been mentioned as a feature in CF [13]. If peroxisomal β -oxidation is reduced in CF, 24:5n-6 and 24:6n-3 can be expected to appear in the gas chromatogram. However, careful inspection of the chromatograms did not reveal the presence of such peaks. Consequently, the low DHA level without a reactive increase in 22:5n-6 could possibly result from a defective elongation of 22:5n-3 and 22:4n-6. EFA deficiency may contribute to the development of lung disease in CF patients. Studies using EFA deficient rabbits and chickens have suggested that EFA shortage predisposes for lung infection with *S. aureus* and *Ps. aeruginosa*, the two most common microorganisms infecting CF patients [34,35]. EFA shortage also seems to be related to different lung surfactant activity in the neonate [36]. Lawrence and Sorrell [37] showed that dietary 20:5n-3 supplementation resulted in a significant reduction in sputum volume and improvement in FEV₁ values and vital capacity in CF patients. Gibson et al [19], showed a significant positive correlation between pulmonary function and levels of 18:2n-6, 20:4n-6, 22:4n-6, 22:5n-3, 22:6n-3 and total n-3 FA in CF. In a study performed by Thompson [38] the only FA showing a significant correlation with pulmonary function was 22:6n-3. Recently Schwartz [39] reviewed the role of PUFA in lung disease, where he points to several trials dealing with supplementation of n-3 FA suggesting an improvement in lung disease. The existence of a correlation between EFA shortage and decreased lung function in CF patients, suggest that correction of this deficiency and particularly of 22:5n-3 and 22:6n-3 could lead to an improvement of lung function, just as we demonstrated in one of our studies [22]. Several studies showed that dietary supplementation of CF patients with fish oils with high amounts of the n-3 FA 20:5n-3 and 22:6n-3 leads to a significantly better incorporation of these FA into plasma, RBC and platelet phospholipids [40-42]. Suppletion of n-3 FA revealed also a significant reduction of the serum concentration of leukotriene B₄, an inflammatory mediator in CF enhancing the influx of neutrophils into the lung [37, 42].

In conclusion, the present study confirmed the presence of a marginal EFA status in pancreatic insufficient CF patients, particularly in plasma PL. This EFA shortage is, amongst others, thought to be a predisposing factor for lung disease in CF patients. The fact that CF patients who received a PPI in addition to the standard treatment showed a less disturbed EFA status compared to CF patients without PPI, suggests that fat malabsorption might, at least partly, be responsible for the EFA deficiency. Because lansoprazole did not normalize steatorrhea completely, we think that additional nutritional supplements of n-3 FA could be useful in children with CF. A larger, placebo controlled study, evaluating the effect of a PPI and n-3 FA supplementation on the EFA status and lung function in CF patients could solve this question.

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CHAPTER EIGHT

Are children with Cystic Fibrosis treated with a proton pump inhibitor at risk for vitamin B12 deficiency?

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8.1. ABSTRACT

Background: In a recent study, the authors demonstrated the beneficial effect of proton-pump inhibitors (PPI) on fat malabsorption and bone mineral content in children with cystic fibrosis (CF). Prolonged use of PPI could result in vitamin B₁₂ deficiency as a consequence of impaired release of vitamin B₁₂ from food in a nonacid environment. The aim of this study was to evaluate the vitamin B₁₂ status of CF patients either treated with a PPI or not by measuring vitamin B₁₂ and homocysteine blood levels, the latter being a sensitive indicator of vitamin B₁₂ deficiency.

Methods: The study population consisted of 20 CF patients, 11 patients treated with a PPI for at least 2 years and 9 CF patients not treated with a PPI, and 10 healthy, age matched control participants. Homocysteine blood levels were measured by high-performance liquid chromatography, and vitamin B₁₂ levels were measured by a competitive protein-binding assay.

Results: Vitamin B₁₂ levels were significantly higher in both CF groups compared with the control participants (PPI+, $P = 0.02$; PPI-, $P = 0.009$). There was no significant difference in vitamin B₁₂ levels between both CF groups. Homocysteine levels were normal and similar in all groups.

Conclusions: Cystic Fibrosis patients treated with a PPI for at least 2 years show no signs of vitamin B₁₂ deficiency.

8.2. INTRODUCTION

Binding to gastric intrinsic factor is the central event in the absorption of vitamin B₁₂. Several steps, however, must occur before this binding can take place. First, food-bound vitamin B₁₂ must be released, mediated by pepsin and acid in the stomach, after which vitamin B₁₂ binds to salivary haptocorrin. At the acid gastric pH environment, the affinity of vitamin B₁₂ for haptocorrin is much higher than for intrinsic factor. Second, in the more alkaline intestinal pH environment, pancreatic enzymes degrade the haptocorrin vitamin B₁₂ complex, and vitamin B₁₂ is transferred to intrinsic factor. Finally, the intrinsic factor-vitamin B₁₂ complex is absorbed in the terminal ileum by phagocytosis after attachment to specific membrane receptors. After absorption vitamin B₁₂ is bound to transcobalamin II in the circulation and subsequently transported intracellularly. Only 6% to 20% of plasma vitamin B₁₂ are attached to the transport protein transcobalamin II. This is the metabolically active fraction that ensures the internalization of vitamin B₁₂ into the body cells, where it acts as a cofactor for the enzymes methionine synthase and methylmalonyl-CoA mutase [1]. Impairment of these enzymatic reactions can result in hyperhomocysteinemia and increased methylmalonic acid blood levels, and clinical manifestations such as anemia and neuropathy [2,3]. The vitamin B₁₂ malabsorption seen in pancreatic insufficient cystic fibrosis (CF) patients has been suggested to be the result of the trypsin-related impaired release of vitamin B₁₂ from haptocorrin. Lindemans et al. [4] showed that the addition of trypsin resulted in an improvement of vitamin B₁₂ absorption in most, but not all, CF patients.

In a recent study, we demonstrated that the use of a proton-pump inhibitor (PPI) in children with CF decreases fat malabsorption and increases body fat and bone mineral content [5]. We were concerned that long-term use of PPIs could result in vitamin B₁₂ deficiency as a result of the impaired release of vitamin B₁₂ from food in a nonacid gastric environment. The purpose of this study was to evaluate the vitamin B₁₂ status of our CF patients by both vitamin B₁₂ and homocysteine blood levels, the latter being a sensitive indicator of vitamin B₁₂ deficiency [3].

8.3. PATIENTS AND METHODS

Twenty CF out-patients from the University Hospital Maastricht were recruited. All children were known to have pancreatic insufficiency on the basis of steatorrhea associated with low fecal chymotrypsin activity. The control group consisted of children admitted to our hospital for minor surgical interventions, for example, orchidopexy and hypospadias. The study was approved by the Ethical Committee of our hospital, and informed patient and parental consent were obtained.

We measured vitamin B₁₂ and homocysteine blood levels in 11 CF patients (mean age, 11.7 years; range, 5.5 - 17.4 years) treated with a PPI (PPI+), in 9 CF patients (mean age, 8.1 years; range, 1.8 - 12.3 years) without a PPI (PPI-), and in 10 healthy, age-matched controls (mean age, 9.2 years; range, 3.3 - 16.2 years). The prescribed

PPI was lansoprazole in a dosage of 15 mg/day orally for at least 2 years. Total homocysteine concentration in plasma (tHcy) was measured by high-performance liquid chromatography [6] and vitamin B₁₂ by a competitive protein binding assay (Perkin-Elmer, Turku, Finland).

8.4 STATISTICS

All data were analyzed using the SPSS statistic program (version 9.0, SPSS Inc., Macintosh, Chicago, IL). Data are presented as mean and range. Vitamin B₁₂ and homocysteine were compared between groups, making use of the Mann-Whitney U test for two independent samples. The Non-parametric Spearman correlation coefficient was used to study the relationship between age, vitamin B₁₂ and tHcy blood levels.

8.5. RESULTS

Mean (and range) of vitamin B₁₂ levels were 553 pmol/L (303 - 907 pmol/L), 637 pmol/L (283 - 1200 pmol/L) and 333 pmol/L (179 - 566 pmol/L) in CF patients receiving a PPI, CF patients without a PPI, and control participants, respectively. In the same patient groups mean and range of tHcy were 7.49 μ mol/L (3.8 - 13.7 mol/L), 6.52 μ mol/L (3.6 - 11.5 mol/L) and 7.76 μ mol/L (4.0 - 10.5 mol/L). Our refe-

FIGURE 8.1. Box plots of vitamin B₁₂ concentrations (pmol/L) in 10 healthy control participants, 9 CF patients not receiving a proton-pump inhibitor (PPI-) and 11 CF patients receiving a PPI (PPI+). The box plot shows the median (central horizontal line), the 25th percentile (lower box border), the 75th percentile (upper box border), and the range of values. The dashed line represents the vitamin B₁₂ upper limit of normal.

FIGURE 8.2. Correlation between homocysteine and vitamin B₁₂ blood levels in healthy control participants ($r = -0.915$; $P < 0.001$).

FIGURE 8.3. Relationship between homocysteine and vitamin B₁₂ plasma levels in CF patients.

rence range of plasma tHcy in 61 children (0 – 10 years of age) was 2.9 – 10.2 $\mu\text{mol/L}$ (mean $\pm$ 2 standard deviations, $6.8 \pm 1.8 \mu\text{mol/L}$). Figure 8.1 shows a box plot of vitamin B₁₂ blood levels in our patient groups. Compared with values obtained in healthy control participants, vitamin B₁₂ levels were significantly higher in CF patients treated with a PPI ($P = 0.02$) and in CF patients without a PPI ($P = 0.008$). There was no

significant difference in vitamin B₁₂ between both CF patients groups ($P = 0.621$). Vitamin B₁₂ blood levels were more than the upper limit of normal (> 630 pmol/L) in 9 of 20 CF patients (5 with and 4 without a PPI). Homocysteine levels were normal and similar in all groups. We found no correlation between age and either vitamin B₁₂ or tHcy levels in our CF patient groups. However, in our control group, a significant positive correlation ($r = 0.78$; $P = 0.008$) was found between age and tHcy levels and a negative correlation ($r = -0.78$; $P = 0.008$) between age and vitamin B₁₂ levels. A strong negative correlation ($r = -0.915$; $P < 0.001$) was found between tHcy and vitamin B₁₂ levels in control participants (figure 8.2.), but not in CF patients (figure 8.3.).

8.6. DISCUSSION

Proton-pump inhibitors, such as lansoprazole, suppress gastric acid secretion and impair the absorption of food cobalamin [7,8]. Several years of ongoing lansoprazole therapy could result in vitamin B₁₂ deficiency. The treatment of our CF patients with a PPI resulted in decreased fat malabsorption and increased body fat and bone mineral content [5]. However we were, concerned that vitamin B₁₂ deficiency could occur in these patients with pancreatic insufficiency receiving long-term treatment with a PPI.

Little is known about the malabsorption of vitamin B₁₂ in children with CF. The malabsorption of crystalline vitamin B₁₂ observed in CF is the result of a mechanism independent of either intrinsic factor secretion or intrinsic factor biological activity [9]. The absorption of crystalline vitamin B₁₂ in CF is corrected in most patients by administration of pancreatic supplements [4,10,11].

We evaluated the effect of prolonged use of a PPI on vitamin B₁₂ and homocysteine blood levels. Our study shows normal homocysteine levels in all groups. Our results are within the reference intervals for children as described in two recent studies [12,13]. Because elevated homocysteine is a sensitive parameter for vitamin B₁₂ deficiency, these results probably rule out vitamin B₁₂ deficiency in our CF patients. As far as we know homocysteine blood levels in CF patients have not been reported previously. In contrast to homocysteine and compared with our normal control participants, vitamin B₁₂ was even found to be elevated in both our CF patient groups regardless of the use of a PPI. We must conclude that the use of a PPI for at least 2 years had no effect on vitamin B₁₂ blood levels in both CF groups. A similar increase in serum vitamin B₁₂ has been reported previously. Lindemans et al. [14] found that in a large group of treated CF patients, serum vitamin B₁₂ levels in general were elevated, despite impaired vitamin B₁₂ absorption. The elevation of vitamin B₁₂ levels in these CF patients occurred while they were not treated with a PPI and was associated with increased levels of transcobalamin II and haptocorrin. Increased levels of transcobalamins have been described in leucocytosis and myeloid leukemia, liver disease, polycythemia vera and autoimmune disorders [15-19]. In patients with alcoholic liver disease, highly elevated vitamin B₁₂ plasma levels were found as a result to leakage of vitamin B₁₂ from

liver tissue into plasma. This resulted in lowered liver tissue vitamin B₁₂ content, increased plasma holo transcobalamin I and III concentration and decreased plasma transcobalamin II [16]. The biological active form of vitamin B₁₂ was significantly depressed because the percentage of vitamin B₁₂ bound to transcobalamin II was diminished [17]. We found a strong negative correlation between homocysteine and vitamin B₁₂ blood levels in our healthy control participants, but not in CF patients. A weaker, but significant, correlation was recently described in healthy Belgian school-children [12]. Because methylcobalamin is a cofactor of homocysteine methyltransferase, such a negative correlation between vitamin B₁₂ and homocysteine blood levels is to be expected. The lack of such a correlation in our CF patients probably implies that because of disturbances in the various forms of circulating cobalamines, the measured 'total' cobalamin blood levels may not be representative of biologically 'active' vitamin B₁₂. Further studies of the different circulating forms of protein bound vitamin B₁₂ and their relationship with plasma homocysteine blood levels are necessary to answer this problem definitively.

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CHAPTER NINE

General conclusions

Han Hendriks

9.1. CONCLUSIONS

Cystic Fibrosis (CF) is relatively rare and the course of the disease is so severe that most patients die at an age of approximately 30 years. All over the world groups of researchers are trying to find out new evidence as to the cause, origin, patho-physiology, course, complications and/or treatment of this life threatening disease. Children with CF together with their parents look anxiously for new ways of increasing the duration and the quality of their lives. As we previously showed a short term beneficial effect of a proton-pump-inhibitor (PPI) in children with CF, we started a new set of studies.

Due to the character of the disease and the restricted number of subjects, the aims of our studies had to be rather limited and consisted of investigating whether or not the pharmacological intervention with a PPI, lansoprazole (Prezal[®]), revealed predictable effects of this medication.

Based on our previous study a decrease of fecal fat excretion and an improvement of body composition could be expected. Possible additional effects of this adjuvant treatment are a decrease of intestinal permeability (IP), an improvement of lung function and of essential fatty acid (EFA) status. As the use of PPIs for a prolonged period might also induce side effects, such as vitamin B₁₂ deficiency, we also looked for evidence of vitamin B₁₂ deficiency in our treated patients. Besides direct evaluation of these effects our study was also intended to provide data, allowing us to perform a power analysis for the planning of further, placebo-controlled, studies. For these purposes an observational study, in 14 children diagnosed with CF and pancreatic insufficiency, was performed, with the approval of the parents, patients and the medical ethical committee.

To maximize the chance to demonstrate an effect, we compared the mean of the whole intervention period to the pre-trial (baseline) values. By this procedure we could demonstrate several effects. As in our previous study we found a decrease of fecal fat excretion immediately after starting the intervention. We also could show that total body fat increased, without a change in fat free mass. A new finding was that bone mineral status improved, probably because of the slow bone turn-over process this was only detected after treatment for 12 months. The effects on body composition, however, turned out to be temporary, as the results at 3 years dropped back to pre-study values. Further we could demonstrate an effect on IP, which in most CF patients decreased, during the first treatment year. We also confirmed the presence of hyperinflation in children with CF, which diminished during therapy. The coincidence between an improvement in fat excretion and body composition on the one hand and a decrease in hyperinflation on the other hand is remarkable, although a causal relationship remains unproven. The effects on EFA are less clear, as only docosa hexaenoic acid (DHA) levels improved significantly. Lastly we demonstrated that after use of a PPI for 3 years neither evidence of vitamin B₁₂ deficiency nor clinical complaints were found. When considering the effects of lansoprazole in CF patients, the following trends were observed. Some effects appeared at once and persisted at the same level during the whole first study year, like the decrease of fecal fat excretion and the hyperinflation

parameters. Other possible effects of PPI such as those on mucosal inflammation (NO_x) and bronchial lability (PEF variables) were more variable and uncertain.

One has to remember, that our main aim was to detect the presence of predictable effects of PPI in a small group of CF patients. We therefore abstain from making any hard recommendations for clinical use of PPIs in children with CF. However, the trends detected in the present study supplies sufficient data to perform a power analysis for future trials.

9.2. POWER ANALYSIS FOR FURTHER STUDIES WITH PPI IN CF

To elucidate definitively the role of PPI in improving fecal fat excretion, body composition, lung hyperinflation, IP, EFA deficiency and other abnormalities in CF, a larger double-blinded, placebo-controlled study would be necessary. This study should be performed in CF children, with comparable stages of the disease. In order to avoid differences in severity of the disease between groups, stratification would be necessary for CF disease score (Schwachman), degree of malnutrition, inflammation and/or lung function. Patients should be randomized in two arms, one PPI group and a placebo group, for one year.

While a cross-over design might be preferable, this would require for several research questions at least a study time of two years. As we do not know how long the carry-over effect is, the total study time would be even longer, up to three years. During such a long period various confounders could occur, interfering with the evaluation of the drug under investigation.

Moreover this study should also assess health economic issues, like cost-effectiveness of the addition of a PPI compared to the standard regime and differences in number of healthy days and number of exacerbations during the intervention. In relatively healthy young CF patients, this would require a large number of subjects. Taking account of our results and using the mean and SD, as computed from the differences between the start (st) and end (e) of study values of each patient variable, we performed a power analysis for future prospective studies. In order to test 2-sided, with $\alpha=0.05$ and $1-\beta=0.8$, the following numbers would be necessary per arm (table 9.1.):

In formula:

$$n_1 = n_2 \geq 2 \times (Z_{\text{sig}} + Z_{\beta})^2 \times \frac{\sigma_{(\text{st-e})}^2}{\mu_{(\text{st-e})}^2} \times \frac{Z_{\text{sig}}^2}{4} \quad (\text{where } z_{\text{sig}} = 1.96 \text{ and } z_{\beta} = 0.842)$$

If we accept that the assessed mean and SD of the changes for each of the variables are sufficient, to prevent a type II error, over 100 patients would have to be included for most variables. In order to resolve all our questions over 100 patients would be needed in each arm. As it would be difficult to find enough children with comparable disease severity (as mentioned above), a multi-center study would certainly be necessary. In the Netherlands just over 400 children with CF are between 5 and 16 years of age. As we demonstrated persisting steatorrhea despite adequate pancreatic enzyme suppletion

TABLE 9.1. Estimated numbers of subjects needed per arm for future studies.
(if $\alpha=0.05$; $1-\beta = 0.8$ and 2-sided testing; based upon values from the present study)

Item		μ_{st-e}	σ_{st-e}	N
Fecal fat excretion: acid steatocrit	(%)	8	14	50
Height	(SDS)	0.1	0.23	85
Weight	(SDS)	0.2	0.40	65
Body composition: TBF	(SDS)	0.3	0.42	32
BMC	(SDS)	0.16	0.21	29
BMD	(SDS)	0.3	0.34	22
Lung Function: TLC	(% predicted)	8	15	57
RV	(% predicted)	35	76	76
RV/TLC	(ratio)	0.05	0.13	108
EFA: DHA	(mg/L)	8	8	17
IP: lactulose recovery	(%)	1.1	2.6	90
L/M ratio	(ratio)	0.1	0.21	71
L/Rh ratio	(ratio)	0.12	0.18	37

For abbreviations see list of abbreviations (page 6).

in only 25% of our patients, this means that in the Netherlands a maximum of 100 patients could be recruited. As the usefulness of a PPI is not the only single question in CF studies and cooperation of so many patients is needed, the involvement of the national and perhaps even the European CF Foundations would be required. Therefore it would be worthwhile, similarly to the CF research consortia in the USA, to organize and establish European CF research consortia.

9.3. PLANS FOR STUDIES IN CF IN MAASTRICHT

Although many studies in the past two decades show some relationship between nutritional status, body composition, lung and gut functioning, the mechanisms involved are still poorly understood. Chronic inflammation in CF can possibly explain several of these disturbances and the effects of a PPI could be related to decreased inflammation. Furthermore, adjuvant therapy with a PPI does not normalize fat absorption completely in all patients; therefore other interventions will still be necessary.

In order to contribute to the study of fat malabsorption in CF patients we are planning the following studies, some in collaboration with the CF center in Rotterdam:

The effects of PPIs on intestinal absorption of fat-soluble vitamins D, K, and bone turnover markers, in order to assess the impact of CF on bone metabolism. We have collected data on this subject (to be submitted, see chapter 4). Based on these results we are

planning to assess the effects of suppletion of vitamin K on bone markers in children and adults with CF.

In order to study inflammation in CF, markers, particularly of pulmonary origin and obtained non-invasively, should become available. Measurements of inflammatory proteins and inflammation-related products in induced sputum and condensated exhaled air are two possibilities to assess pulmonary inflammation in a much less invasive way than Broncho-Alveolar-Lavage. These types of inflammation markers will be evaluated in relation to systemic inflammation, lung function parameters and the nutritional status.

Evaluation of the effects of fish-oil suppletion, a combination of several (n-3)-EFAs, DHA enriched diet, in combination with a PPI on the EFA status and inflammatory parameters in CF patients.

CHAPTER TEN

Summary in English and Dutch

Han Hendriks

SUMMARY OF THE DIFFERENT CHAPTERS

In chapter one the rationale for the use of a proton pump inhibitor (PPI) in Cystic Fibrosis (CF) patients is described. Persisting malabsorption is still a problem in many CF patients, despite supplementation with pancreatic enzymes. Factors involved in malabsorption, as reported in the literature, are given and indicate the importance of duodenal acidity. Normal gastric acid production is buffered with pancreatic bicarbonate, which production is hampered early and severely in CF patients. By adding a proton pump inhibitor (PPI) to the standard regime of CF children with ongoing steatorrhea, despite supplementation with pancreatic enzymes, gastric acid production is reduced which will allow duodenal pH to rise, resulting in a more efficient utilization of pancreatic enzyme capsules and better fat absorption.

Chapter two describes the aims, the patients and methods of the complete study protocol. Our previous, short-term, study had demonstrated a possible effect of administration of PPI on fecal fat excretion and total body fat (TBF) of CF patients. A new observational study was planned, to estimate these as well as additional effects of lansoprazole treatment administered during one year, on several biological variables in children with CF. This study was primarily aimed at looking for possible predictable effects of PPI in children with CF. As it was possible to include only a limited number of patients no definitive conclusions or recommendations as to the clinical use of PPI in CF can be made.

In chapter three we report growth data of our young CF patients. In contrast to most published studies, reporting cross-sectional growth data in large CF populations, we used longitudinal growth data to assess growth patterns. We demonstrated that stunting contributed more to growth retardation than wasting, and therefore the use of weight-for-age growth charts instead of weight-for-height reference values to assess growth failure in a CF patient will be more appropriate. These data also support the use of for screening programs for CF, as growth was severely retarded in 80% of our patients at initial assessment and early intervention has been shown to prevent stunting in these patients.

Chapter four described the evaluation of both the short term (one year) and long term (3 years) effects of adjuvant therapy with a PPI on body composition in CF. Fecal fat excretion was monitored throughout the year by the acid steatocrit method. Body composition was assessed at the start of the study and at 6, 12 and 36 months by dual energy X-ray absorptiometry (DEXA). Body composition data are expressed in standard deviation scores (SDS) instead of absolute values, in order to take into account the normal growth process in children. The addition of a PPI to the treatment of young CF patients resulted in an immediate and long lasting decrease of fat excretion. During the first year of the intervention an improvement not only of TBF, but also of bone mineral content (BMC) and bone mineral density (BMD) occurred. As body composition values at 3 years returned to pre-treatment values, our results demonstrated the beneficial

effect of PPI on body composition in CF to be temporary. We propose to assess body composition by DEXA in CF patients regularly, allowing a better monitoring of the nutritional condition compared to anthropometric data alone.

The changes in lung function parameters in CF patients, during the use of PPIs, were described in chapter five. Our results confirmed the presence of hyperinflation in CF and showed a decrease of hyperinflation during the use of a PPI, which can only partly explain the improvement of inspiratory muscle performance. Whether the association between an improvement of body composition and a decrease of lung hyperinflation is causal or coincidental cannot be ascertained from our study. Furthermore daily recording of peak expiratory flow (PEF) confirmed that PEF variability is increased in CF, which might point to the presence of bronchial lability in these patients.

In chapter six we confirmed the well-known and severely increased intestinal permeability (IP) in CF patients. Our results showed that the increased IP, at least partly, can be corrected by the use of a PPI during one year. As the paracellular permeability pathway mainly improved, we suggested that acid luminal contents might damage the intestinal mucosa, particularly the tight junctions, contributing to the increase of IP in CF.

Biochemical results concerning signs of essential fatty acid (EFA) deficiency in our patients were presented in chapter seven and showed plasma phospholipid fatty acids to be mainly affected. Although the EFA status of patients receiving PPIs showed some trend towards normalization, the only difference between CF patients with PPI and those without, consisted in a higher plasma docosa hexaenoic acid (DHA) level in the PPI group. DHA has been reported to play a role in CF pathophysiology and CFTR-related disease. These results suggest that fat malabsorption contributes to the EFA deficiency of CF patients.

As explained in chapter one, most adverse effects of PPI are mild and probably do not outweigh the beneficial effects. However, prolonged use of a PPI may result in vitamin B₁₂ deficiency due to impaired release of vitamin B₁₂ from food in a non-acid environment.

In chapter eight we showed that vitamin B₁₂ is significantly higher in CF patients compared to healthy controls, independent of PPI administration. In PPI treated and non-PPI treated CF patients homocysteine blood levels, a sensitive indicator of vitamin B₁₂ deficiency, were in the normal range. As there was no significant difference in both vitamin B₁₂ and homocysteine blood levels between both CF groups, we assumed that PPI's, even when used for nearly three years, had no adverse effect on vitamin B₁₂ status.

GENERAL SUMMARY

The effects of decreasing luminal acidity on several biological variables was studied, using a pharmaceutical intervention in CF patients. Our main aim was to look for possible beneficial effects of lansoprazole on fecal fat excretion, body composition, lung function, gut permeability and EFA status without causing important side effects, like a vitamin B₁₂ deficiency. As our aim was mainly to look for predictable effects of PPIs in CF, we chose for an open, observational study in a small patient group which makes it necessary to interpret our results cautiously. Similarly to our findings in a previous study, we confirmed that the use of lansoprazole resulted in a decrease of stool fat and a temporary increase of body fat and bone mineral status. Additional effects showed by the present thesis are a decrease in hyperinflation of the lung, bronchial lability, intestinal permeability and improvement of EFA status without evidence of vitamin B₁₂ deficiency.

Although possibly important changes of clinical variables in children with CF have been observed, their causal relationship with the use of a PPI remains unproven due to the observational design and the small number of patients involved. However our results do now allow us to perform a power analysis in order to estimate the number of patients needed for future double-blind, placebo-controlled studies. Results of that evaluation show that large multi-center (multi-national?) studies would be necessary, involving high costs. This, in addition to the fact that several other questions have to be solved urgently in CF, makes it unlikely that such studies will be performed in the near future.

Although the use of PPIs in the Netherlands is already general practice in adult CF patients with persisting steatorrhea despite adequate supplementation of pancreatic enzymes, hard evidence for the use of this treatment in children is lacking. Although the present thesis suggests possible beneficial effects of adjuvant therapy with a PPI in CF children with ongoing steatorrhea, it does not allow us to recommend the routine clinical use of PPI in these children. Based on our limited data we encourage the early detection of malabsorption and malnutrition in CF. To this end the yearly assessments of body composition by DEXA seems reasonable. If despite adequate pancreatic enzyme supplementation malabsorption and malnutrition persists, a trial with a PPI seems warranted.

SAMENVATTING VAN DE AFZONDERLIJKE HOOFDSTUKKEN.

In hoofdstuk één is de rationale voor het gebruik van protonpomp remmers (PPI) in patiënten met Cystic Fibrosis (CF) beschreven. Ondanks de inname van pancreas enzymen blijft verlies van vet met de ontlasting een probleem voor menig CF patiënt. Middels literatuurstudie werd een overzicht gegeven van de verschillende factoren die betrokken zijn bij de zogenaamde malabsorptie, met daarbij speciale aandacht voor de zuurgraad in het duodenum. Normaal gesproken wordt maagzuur in het duodenum gebufferd door bicarbonaat uit de pancreas. Bij CF patiënten is de productie van bicarbonaat, al in een vroege fase van de ziekte, ernstig aangetast. Door een PPI toe te voegen aan de standaard behandeling van CF patiënten met malabsorptie, ondanks suppletie van pancreas enzymen, wordt minder maagzuur geproduceerd. Hierdoor zal de zuurgraad in het duodenum lager zijn, resulterend in een meer efficiënt gebruik van de pancreas enzymen, zich vooral uitend in een betere vetabsorptie.

Hoofdstuk twee beschreef de doelstellingen, de patiënten en de methoden van het complete studie protocol. In een eerdere, korte termijn, studie hadden wij al een mogelijk effect van toediening van PPI op de fecale vet excretie en totaal lichaamsvet (TBF) in kinderen met CF aannemelijk gemaakt. Een nieuwe, observationele studie, werd gepland om deze én additionele effecten van lansoprazol (een PPI) toediening gedurende een jaar, op verschillende biologische variabelen in kinderen met CF te meten. Deze studie was primair bedoeld om te onderzoeken of er voorspelbare effecten van PPI in kinderen met CF zijn en hoe groot deze effecten dan zijn. Omdat slechts een beperkt aantal patiënten in deze studie bestudeerd is, kunnen derhalve geen definitieve conclusies of aanbevelingen aangaande het klinisch gebruik van PPI in kinderen met CF gemaakt worden.

In hoofdstuk drie rapporteerden wij de groeigegevens van onze jonge CF patiënten. In tegenstelling tot de meeste gepubliceerde studies, met cross-sectionele groeidata in grote CF populaties, gebruikten wij longitudinale groeigegevens om groeipatronen vast te stellen. Wij toonden aan dat achterblijven in lengtegroei (stunting) meer bijdraagt aan de groeivertraging dan te weinig aankomen in gewicht naar lengte alleen (wasting). Hierbij werden de CF kinderen vergeleken met naar leeftijd gematchte, gezonde kinderen. Om een groeivertraging te meten is daarom het gebruik van gewicht naar leeftijd groeicurven adequater dan het gebruik van gewicht naar lengte groeicurven. Verder ondersteunen deze bevindingen de noodzaak voor neonatale screening naar CF. Enerzijds omdat reeds bij de eerste meting een groeivertraging in 80% van onze patiënten aanwezig was en anderzijds omdat van vroege interventie is aangetoond dat "stunting" bij deze patiënten wordt voorkomen.

Hoofdstuk vier beschreef zowel de korte termijn (één jaar) als de lange termijn (3 jaren) effecten van aanvullende therapie met een PPI op de lichaamssamenstelling bij CF patiënten. Het verlies van vet met de ontlasting werd gedurende één jaar gemeten met

de “zure steatocrit” methode. De lichaamssamenstelling werd gemeten bij de start van de studie en na 6, 12 en 36 maanden met behulp van dual energy X-ray absorptiometrie (DEXA). De gegevens van de lichaamssamenstelling werden uitgedrukt in standaard deviatie scores (SDS) in plaats van absolute waarden, om rekening te kunnen houden met het normale groeiproces in kinderen. De toevoeging van een PPI aan de behandeling van jonge CF patiënten resulteerde in een onmiddellijke en lang aanhoudende afname van het verlies van vet met de ontlasting. Gedurende het eerste jaar van de interventie trad er niet alleen een verbetering van het TBF op, maar ook van het mineraal gehalte (BMC) en minerale densiteit van bot (BMD). Omdat de waarden van de lichaamssamenstelling na 3 jaren gelijk waren aan de waarden bij de start, toonden onze resultaten slechts een tijdelijk positief effect van PPI op de lichaamssamenstelling in kinderen met CF. Wij stellen voor om regelmatig de lichaamssamenstelling te meten van CF patiënten, met behulp van DEXA, waardoor een betere monitoring van de voedingstoestand mogelijk is, dan alleen met wegen en meten.

In hoofdstuk vijf werden de veranderingen in longfunctieparameters tijdens het gebruik van PPI's in onze CF patiënten beschreven. De resultaten bevestigden de aanwezigheid van een hyperinflatie van de longen in CF patiënten en toonden, tijdens het gebruik van een PPI, een afname van hyperinflatie aan. Dit kon slechts gedeeltelijk de verbetering van de kracht van de inademiings-spiers verklaren. Naar aanleiding van deze studie kon niet bepaald worden of de associatie tussen een verbeterde lichaamssamenstelling en een afname van de longhyperinflatie causaal dan wel toeval was. Tevens bevestigden dagelijkse registratie van uitademings-piekstroom (PEF), dat de variabiliteit in PEF toegenomen is in CF patiënten, wat zou kunnen duiden op de aanwezigheid van bronchiale labiliteit.

In hoofdstuk zes bevestigden wij de bekende, fors verhoogde doorlaatbaarheid van de darmen in CF patiënten, ook wel intestinale permeabiliteit (IP) genoemd. Onze resultaten toonden aan dat de toegenomen IP tenminste deels gecorrigeerd werden door het gebruik van een PPI gedurende een jaar. Omdat voornamelijk de permeabiliteit tussen darmcellen (paracellulaire route) verbeterde, nemen wij aan dat de zure darminhoud in CF patiënten vooral de bindingen tussen darmcellen beïnvloeden met een toegenomen IP als gevolg.

De biochemische bevindingen, voor aanwijzingen van een tekort aan essentiële vetzuren (EFA), in onze CF patiënten werden gepresenteerd in hoofdstuk zeven. Het tekort bleek met name in de vetzuren van plasma fosfolipiden. Hoewel de meeste EFA van patiënten, die PPI's gebruikten, een geringe trend naar normalisatie toonden, bleek alleen het verschil in plasma docosa hexaenoic acid (DHA) in de PPI-groep significant hoger dan in de niet gebruikers groep. Van DHA is beschreven dat het een rol speelt in de pathofysiologie van CF en CFTR-gerelateerde aandoeningen. Onze resultaten suggereren dat vetmalabsorptie bijdraagt aan de EFA deficiëntie in CF patiënten.

Zoals beschreven in hoofdstuk één, zijn de meeste bijwerkingen van PPI's mild en wegen waarschijnlijk niet op tegen de gunstige effecten. Echter, langdurig gebruik van een PPI zou kunnen leiden tot een vitamine B₁₂ tekort. Dit is te verwachten omdat aan voedsel gebonden vitamine B₁₂ in een niet-zure omgeving minder loslaat en daardoor niet opgenomen wordt.

In hoofdstuk acht tonen wij juist aan dat, in vergelijking met gezonde controle patiënten, plasma vitamine B₁₂ significant hoger is in CF patiënten, onafhankelijk van het gebruik van een PPI. Zowel in de PPI behandelde groep, als de niet-PPI behandelde groep zijn de homocysteïne serumwaarden, een sensitieve indicator van vitamine B₁₂ tekort, in de normale range. Omdat er geen significant verschil was in zowel vitamin B₁₂ als homocysteïne serumwaarden tussen beide CF groepen, concludeerden wij dat PPI's, zelfs indien gedurende bijna drie jaar gebruikt, geen invloed hadden op de vitamine B₁₂ status.

ALGEMENE SAMENVATTING

De effecten van een afgenomen zuurgraad van de darminhoud op diverse biologische variabelen is bestudeerd, middels een medicamenteuze interventie bij kinderen met CF. Het hoofddoel was om mogelijke gunstige effecten van lansoprazole, een PPI, aan te tonen op vetverlies met de ontlasting, lichaamssamenstelling, longfunctie, darm permeabiliteit en EFA status bij kindere met CF. Tevens werd gekeken naar mogelijke bijwerkingen van langdurig PPI gebruik, zoals een vitamine B₁₂ tekort. Omdat de doelstellingen vooral gericht waren op het aantonen van specifieke effecten van PPI's in CF, was het mogelijk om te kiezen voor een open, observationele studie in een patiënten groep met kleine aantallen. Dit maakt het echter nodig om onze resultaten voorzichtig te interpreteren en zijn de resultaten derhalve niet conclusief. De bevindingen uit onze eerdere studie, werden bevestigd, het gebruik van lansoprazole resulteerde in een afname van vet in de ontlasting met, een tijdelijke, toename van totaal lichaamsvet en botmineralisatie. Andere uitkomsten van onze studie zijn zowel een afname van de hyperinflatie van de long, bronchiale labiliteit en darm permeabiliteit, als een verbetering van de EFA status en dat zonder een vitamine B₁₂ tekort te veroorzaken.

Hoewel belangrijke veranderingen in klinische variabelen in kinderen met CF geobserveerd zijn, is een oorzakelijke relatie met het gebruik van een PPI niet bewezen als gevolg van het observationele design en het kleine aantal patiënten. Echter met de verkregen resultaten hebben we een power-analyse uitgevoerd, om het benodigde aantal patiënten voor een toekomstige dubbel-blinde, placebo-gecontroleerde studie te berekenen. Resultaten van deze evaluatie tonen dat grote multi-center (multi-nationale?) studies nodig zijn, leidend tot hoge kosten. Dit naast het feit dat andere, meer dringende, vraagstellingen opgelost dienen te worden in CF, maakt het

onwaarschijnlijk dat dergelijke studies in de nabije toekomst zullen worden uitgevoerd.

Het gebruik van PPI in Nederland is algemeen gebruik bij volwassen CF patiënten, die vetverlies via de ontlasting houden ondanks gebruik van pancreasenzymen. Daarentegen bestaat er geen bewijs voor een dergelijke behandeling bij kinderen. Het onderzoek in dit proefschrift suggereert mogelijke gunstige effecten van toevoegen van een PPI in kinderen met CF met voortgaand vetverlies met de ontlasting, maar het staat ons niet toe om het routinematige gebruik van PPI in deze kinderen aan te bevelen. Gebaseerd op onze beperkte gegevens moedigen wij de vroege detectie van malabsorptie en ondervoeding bij patiënten met CF aan. Met dit als doel lijkt het jaarlijks meten van de lichaamssamenstelling met DEXA logisch. Als ondanks adequate hoeveelheden pancreasenzymen, malabsorptie en ondervoeding persisteren, lijkt toevoegen van een PPI verantwoord.

CHAPTER ELEVEN

Dankwoord en Curriculum Vitae

Han Hendriks

DANKWOORD

Als er één waarheid is over het schrijven van een proefschrift dan is het wel dat het niet het werk is van slechts één persoon. Ik wil dan ook bij voorbaat iedereen zeer hartelijk danken voor hun bijdrage aan het tot stand komen van dit proefschrift, ook degenen die ik nu niet vernoem. Elke hulp, hoe klein ook, is van belang en wordt zeker ook door mij niet vergeten. Maar een aantal personen verdienen speciale dank en die wil ik nu verwoorden.

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aandacht uitgaat naar grote, gerandomiseerde, dubbel blinde en placebo gecontroleerde studies, moeten wij niet vergeten dat studies, zoals in dit proefschrift beschreven, mede de basis vormen waarop dergelijke studies hun aantallen en design baseren. Verder dient bij een relatief zeldzame ziekte, zoals Cystic Fibrosis, er zoveel tegelijkertijd onderzocht te worden dat dergelijke grote en dure studies nooit uitgevoerd worden. Daarentegen is de noodzaak groot om met beide handen iedere verbetering van het beloop van deze gevreesde aandoening aan te pakken, opdat patiënten niet alleen langer leven maar ook hun kwaliteit van leven verbeterd.

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CURRICULUM VITAE

De schrijver van dit boek werd op 19 maart 1958 geboren, in Maastricht aan de voet van de Sint Pietersberg. In 1976 behaalde hij het VWO-diploma, aan het toenmalige Henric van Veldeke College te Maastricht. Nog hetzelfde jaar startte hij met zijn studie Geneeskunde aan de Katholieke Universiteit van Nijmegen, alwaar hij op 14 oktober 1983 zijn artsexamen behaalde. Zijn militaire dienstplicht, als reserve kapitein arts bij de stafcie van de 4e divisie, mocht hij een maand eerder beëindigen, om zijn opleiding tot kinderarts te starten. Van februari 1985 tot mei 1988 werd hij opgeleid in het St. Radboud ziekenhuis te Nijmegen (opleider: prof.dr. GBA Stoelinga) om daarna tot februari 1989 zijn opleiding voort te zetten in het Academisch Ziekenhuis Maastricht (opleider: prof.dr. RH Kuyten).

Sinds 1 februari 1989 is hij werkzaam als kinderarts bij de vakgroep Kindergeneeskunde van het AZM. Na een periode gefunctioneerd te hebben als chef de clinique, ging hij zich verder specialiseren in de kinderlongziekten. Daartoe werkte hij van mei – december 1990 in het Academisch Ziekenhuis Groningen (opleider: prof.dr. K.Knol), verdere stages werden gevolgd in het AZM bij prof.dr. EFM Wouters en in Boedapest, Hongarije, bij dr. E. Szekely.

Sinds januari 1991 werkt hij als kinderarts op het gebied van de kinderlongziekten, daarbij krijgt hij sinds januari 1998 versterking van Rijn Jöbssis en sinds april 2000 van Edward Dompeling. Na invoering van een registratie van subspecialismen binnen de Kindergeneeskunde, werd hij in januari 1999 retrograad geregistreerd als kinderarts-pulmonoloog. Momenteel bekleedt hij meerdere functies binnen het AZM, het coördinatorschap van het CF-team en de geneesmiddelen commissie. Ook binnen de vakgroep Kindergeneeskunde worden diverse functies bekleed binnen de refereer, de opleidings en de status commissie. Tevens is hij sinds 1996 hoofd van de polikliniek Kindergeneeskunde en lid van het dagelijks bestuur van de vakgroep Kindergeneeskunde.