BEAUMONT HOSPITAL

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KIDNEY DISEASE

A Guide for Patients



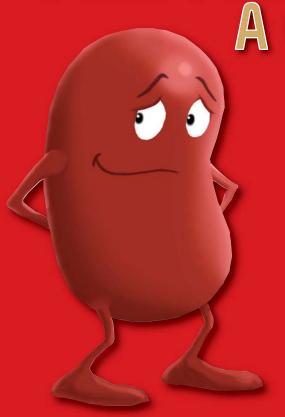
Petrina Donnelly, CNM, Helen Dunne, CNM Prof. Peter J. Conlon, FRCPI

4th Edition

Re-edited by Ruth O'Malley, Olive McEnroe, Binu Vasu









IRISH KIDNEY ASSOCIATION (CLG)

The Irish Kidney Association CLG, Company Limited by Guarantee, is a national voluntary organisation of patients, family carers and supporters which offers support for all patients with end stage kidney disease. Through its 25 local branches, patients can meet other kidney patients and share experiences, problems and, most importantly, solutions.

The Irish Kidney Association Head Office is located in the west of Dublin, named 'Donor House'. The office is the main administrative and services centre for the Association. The staff is made up of Chief Executive, Accountant, Patient Support Office, Office Manager and Personal Assistant to CE, National Projects Manager, Renal Counsellor, Dialysis Holiday Coordinator, Receptionist, IT Developer, Social Media Co-ordinator and the Support Centre Managers. Individual names and email addresses for the staff are available on the IKA website.



The IKA provides holidays every year, for kidney patients who are either receiving dialysis treatment or are transplanted. It also has a Support Centre, in the grounds of Beaumont Hospital, which offers on-campus accommodation for kidney patients and their families attending any Dublin hospital and short-term accommodation for the families of seriously ill patients from outside the Dublin area.

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CHARITY REGISTRATION NUMBER: 20011260



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IKA RENAL SUPPORT CENTRE

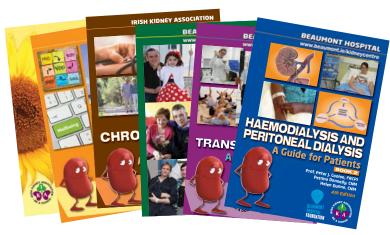
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PREFACE

n 1983, Dr Michael Carmody compiled the first book of Living with Kidney Failure. For more than 20 years, this book was used as the major education booklet for patients experiencing kidney disease. During the last 20 years, the technology concerning the treatment of kidney disease has changed radically. As a result we have produced this series of books to assist patients and their families when diagnosed with kidney disease.

This is the fourth edition of **BOOK 1** in a series of seven books, aimed at helping patients with kidney disease learn more about their illness. Book 1 deals with the functions of the kidney, types of kidney diseases, diagnostic tests and medicines used to treat kidney conditions.



BOOK 2 deals with Haemodialysis and Peritoneal Dialysis; **BOOK 3** - Kidney Transplantation; **BOOK 4** addresses the area of the 'living donor' programme; **BOOK 5** is aimed at patients living with Chronic Kidney Disease; **BOOK 6** - Emotional Wellbeing and **BOOK 7** - deals with the Conservative Care of Kidney Failure.

Please use these books as a guide and reference tool but, any worries or issues you have should be discussed with your medical team. The text includes contributions from many members of the Beaumont Hospital Renal Unit team and has been supported by the Beaumont Transplant Foundation together with the Irish Kidney Association.

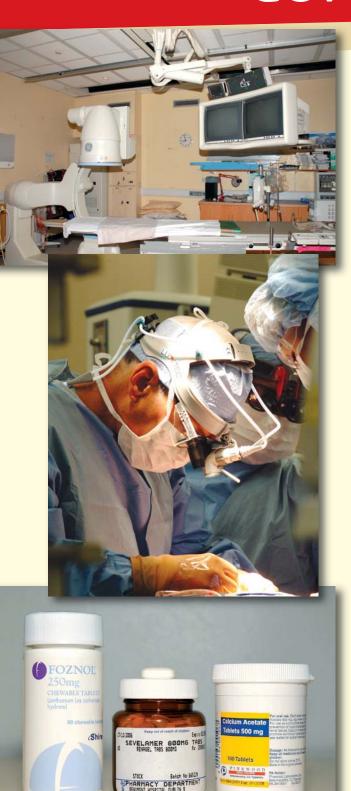
We do hope you find it helpful.

Petrina Donnelly CNM Helen Dunne CNM Prof. Peter J Conlon FRCPI





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The information contained within this book is correct at time of going to press. This book essentially pertains to the practices at Beaumont Hospital. Other Kidney Units may use different practices. This book should be used as a guide and reference tool only.

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The books have been printed and distributed by the Irish Kidney Association. Further copies are available from IKA, Donor House, Parkwest, Dublin D12 P5V6. Ph: 01-6205306.

THE URINARY SYSTEM

CHAPTER 1

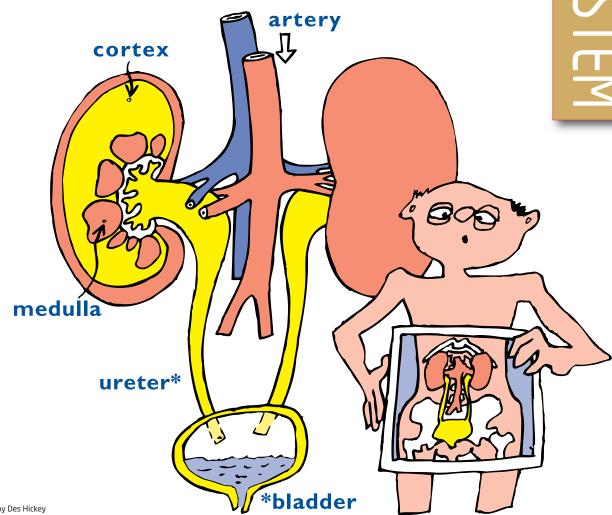
he urinary system is one of the body's systems which helps us to dispose of the waste products naturally produced within the body. The main structures in this system are:

Two kidneys - which lie behind the other major organs in the lower back area. They are bean-shaped organs and measure about 11cm long, 6cm wide and 3cm deep. They have 5 main functions, which will be discussed at a later stage.

Two ureters - (tube-like features) which run from the kidneys to the bladder carrying urine.

One bladder - which collects urine from the kidneys, via the ureters, and stores it temporarily.

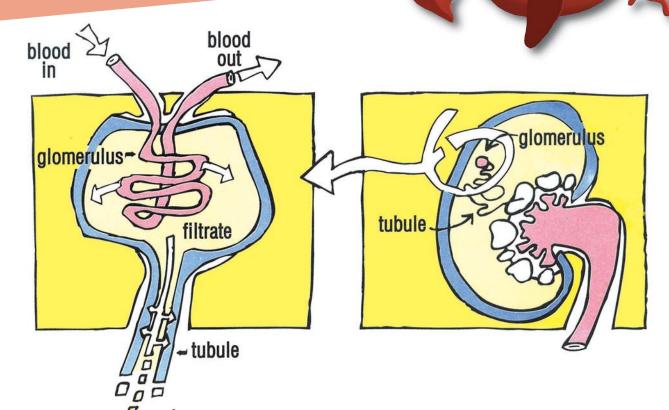
One urethra - through which the urine is excreted out of the body, allowing the bladder to empty and dispose of the waste.



WHAT THE KIDNEYS DO:

The kidneys...

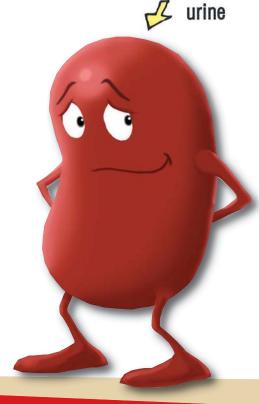
- Filter and remove the waste products of the body.
- Remove excess water from the body.
- Help in the production of red blood cells, which carry oxygen around the body.
- Help maintain healthy bones.
- Help regulate blood pressure in the body.





As blood flows through the body it picks up waste and carries this to the kidneys using the kidney arteries. The waste in your blood comes from the normal breakdown of active tissues and from the food you eat. Your body uses food for energy and self-repair. After the body has taken what it needs, from the food, the waste is sent to the blood. The kidneys filter out the waste products and excess fluids from the body and dispose of them in the form of urine, via the bladder. The clean blood flows back to the other parts of the body. If your kidneys did not remove this waste, it would build up in the blood and cause damage to your body.

The actual filtering occurs in tiny units inside your kidneys called nephrons. Each kidney contains about a million nephrons. In the nephron, a glomerulus (which is a tiny blood vessel or capillary) intertwines with a urine collecting tube called tubules. A complicated chemical exchange takes place, as waste materials and water in your blood enter your urinary system.



SOME EXAMPLES OF WASTE PRODUCTS INCLUDE:

Urea - Blood carries protein from your diet to the cells to fight disease and repair muscle. Whatever protein is not used is put back into the bloodstream in the form of urea for excretion. Too much urea, in the blood, is known as uraemia.

Potassium - A mineral absorbed into the bloodstream from many fruits and vegetables such as oranges, bananas and potatoes. Potassium regulates the heart rate. Healthy kidneys remove excess potassium, from the blood stream, as a waste product.

Creatinine - A waste product in the blood created by the breakdown of muscle cells during activity. The levels vary according to the size of the individual, i.e. the muscle mass of the person.

Sodium - A chemical absorbed in the blood stream from food containing salts. Excess sodium in the blood may cause a rise in blood pressure as it plays a vital role in regulating the amount of fluid in the blood.

Chloride - Like sodium, this chemical helps maintain a balance of fluid in the body. It is also absorbed from salted foods and is absorbed in the bloodstream in the bowel.

In addition to removing waste, the kidneys have other important functions. These are carried out with the help of three hormones, which are released in the kidneys.

Erythropoietin (eh-rith-ro-poy-eh-tin) or Epo is a hormone that is secreted by your kidneys and stimulates the bone marrow to produce red

blood cells, which carry oxygen in the blood, to the cells in the body.

Renin (ren-in) is a hormone that the kidneys secrete to help regulate the blood pressure through a chemical process within the blood stream.

Calcitriol (kal-suh-try-ul) is the active form of Vitamin D that the kidneys

secrete to help maintain healthy bones by maintaining a chemical balance between calcium and phosphate in the blood.

HOW DOES KIDNEY DISEASE AFFECT YOUR BODY?

Kidney disease can affect you in a number of different ways. These include:

- Proteinuria (protein in the urine)
- Haematuria (blood in the urine)
- Hypertension (high blood pressure)
- Elevated Serum Creatinine (kidney disease)
- Kidney Stones
- Recurrent Urine Infections

Proteinuria

"Proteinuria

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Proteinuria or protein in the urine is frequently the earliest symptom of kidney disease. You will have read, in the previous section, how the kidney works and that the kidney has about a million filters. When the kidney is healthy it allows very little protein into the urine. If these filters become leaky, small amounts of protein will leak into the urine. This is frequently an early sign of kidney trouble long before the kidney function itself begins to deteriorate.

Doctors frequently test patient's urine for the presence of blood or protein, to try to detect kidney disease early. There are many causes of protein in the urine, including diabetes and glomerulonephritis. Whilst your doctor will conduct a number of special blood tests, to try to determine

the underlying cause, it may be necessary to undergo a kidney biopsy (see page 26), to establish the exact cause of the protein.

Patients who have very large amounts of protein in the urine, (greater than 3 grams), are described as having nephrotic syndrome. Patients with nephrotic syndrome frequently have swollen legs.

Haematuria

Blood in the urine (haematuria) can either be present in amounts that you can see (macrascopic) or in amounts that you cannot see (microscopic) in which it is only detected with urine testing. Blood in the urine may not appear red but more like strong coloured tea.

Blood in the urine is frequently an alarming symptom and it should never be ignored. However, it only takes a few drops of blood for the urine to turn red. There are a large number of potential causes of blood in the urine including: urine infection, kidney stones, kidney or bladder tumours and inflammation in the kidney called glomerulonephritis (GN).

If you have haematuria, the first thing your doctor will do is to make sure you do not have a urine infection or bladder or kidney tumour. To do this, you will usually need to have a number of scans of the kidney and may well need a cystoscopy. A cystoscopy is a test in which a camera with a light is inserted into the bladder. If these tests are normal your doctor will

then focus on determining if the blood is coming from kidney inflammation or glomerul-onephritis. This may require further specific blood and urine tests or a kidney biopsy (see page 26).

High Blood Pressure

High blood pressure (hypertension) may arise due to, or as a result of, kidney disease. lt is important, however, realise that the vast majority of people, with high blood pressure, have entirely normal kidney function. If you have high blood pressure, it is very important to treat it as it will help preserve your kidney function and reduce the risk of developing a stroke or heart attack.

Elevated Serum Creatinine

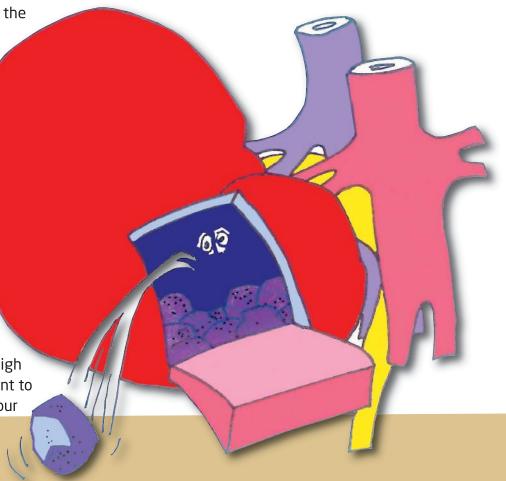
"Blood in the urine is frequently an alarming symptom and it should never be ignored."

The serum creatinine is a blood test that is used to monitor kidney function. The serum creatinine may be elevated as an early sign of kidney disease long before there are any symptoms evident of kidney disease (see Chapter 2). Blood tests, identifying elevated serum creatinine,

are mainly undertaken, as a routine, during an annual physical medical examination. If serum creatinine is elevated, it may be as a result of any of the kidney diseases discussed below.

Kidney Stones

Kidney stones occur when a tiny fragment of crystal develops within the kidney or the tube coming from the kidney, called the ureter. A kidney stone mainly produces very severe colicky pain. The pain of a kidney stone is described as being more severe than that of labour pains. The treatment of a kidney stone will consist of pain medication initially mainly in the form of an injection.



Subsequently, the doctor/urologist may attempt to extract the stone. "Recurrent urine This may occur by way of a number infections are a of methods, depending on the size common kidney and position of the kidney stone. complaint, particularly The urologist may pass a scope amongst into the bladder and try to grasp young women," the stone. They may also try to bypass the stone temporarily with a stent, or may try to dissolve the stone with a machine called a Lithotripsey. Occasionally, the urologist may be required to perform an open operation, on the kidney, in order to surgically remove the stone.

Once the stones are removed, the team will attempt to determine their underlying cause. Conditions associated with recurrent kidney stones include:

- Not drinking enough water. If you have kidney stones, it is necessary to drink 3 litres of water a day.
- High levels of calcium in the urine (hypercalciuria). This condition, in which some people pass too much calcium into the urine, can occur for unknown reasons.
- High levels of calcium in the blood (hypercalcemia). This may occur as a result of a gland in the neck called the parathyroid gland becoming overactive. If this is the case you may need to undergo a small operation to remove part of the gland. It may also occur as a result of consuming too much calcium in the diet.

"If you have kidney stones it is necessary to drink 3 litres of water a day."

High urinary oxalate (oxalosis). Oxalate is a constituent of certain foods. If you are diagnosed as having too much oxalate your doctor may prescribe a diet, high in calcium, as this helps to prevent kidney stones in this situation.

of kidney stones and is inherited (runs in families). This occurs as a result of the body being unable to handle the amino acid cysteine. The treatment recommended for this condition is to consume large amounts of water. In addition, medications are sometimes prescribed to help reduce the frequency of kidney stone formation.

The investigation of patients with kidney stones will consist of a number of 24 hour urine collections and blood tests to determine which of the above

conditions is responsible for the kidney stones. One of the urine collection bottles contains acid and it is important that it is handled with care and kept far away from children.

(See pg 19 for instructions on 24 hour urine collections).



Recurrent urine infections are a common kidney complaint, particularly amongst young women. The vast majority of patients, with recurrent urinary infection, do not have any underlying structural kidney disease. It is important, however, to eliminate kidney disease as a possible cause by way of performing kidney x-rays. Sometimes long-term (6 months) prophylactic antibiotics are used to eliminate any underlying infection.

CHAPTER 2

hronic kidney disease (CKD) is a condition in which the kidneys cannot perform their normal functions. Your kidneys lose the majority of their filtering ability and, as a result, fluid and waste accumulates in your body.

When CKD happens suddenly, it is known as **ACUTE KIDNEY INJURY** (AKI). The most common causes of acute damage to the kidneys are:

- Decreased blood flow to the kidney: this may occur when there is extremely low blood pressure caused by trauma, complicated surgery, septic shock, haemorrhage, burns, associated dehydration or other severe or complicated illnesses.
- Over-exposure to metals, solvents, x-ray dye, certain antibiotics and other medications or substances.
- Acute Tubular Necrosis (ATN) may occur when the tissues are not getting enough oxygen.

Short-term treatment may be needed for acute kidney injury, but the kidneys usually recover on their own. However, if the cause of the acute kidney injury persists, there can be permanent damage to the kidney, which will lead to CKD.

CHRONIC KIDNEY DISEASE (CKD) usually develops slowly, with few signs or symptoms in the early stages. You may still be passing normal amounts of urine, but it will be poor quality, and waste products, which should normally be filtered out, will remain in the body. Many people with CKD do not realise they have a problem until their kidney function has decreased to less than 25 percent of normal (CKD4). This damage usually occurs slowly, and is not reversible.

The rate of deterioration of kidney function is variable, ranging from more than ten years to only a few months.

Eventually, the kidneys can only function at less than 10 percent of normal capacity. The kidneys have almost stopped working at this stage and treatment, in the form of dialysis or a kidney transplant is required to take over the work of the kidneys and maintain life.

This is commonly known as **END STAGE KIDNEY DISEASE** (ESKD). In other words, kidney damage is irreversible and cannot be controlled by conservative management alone. When kidneys reach 'end-stage', they never recover.

PROGRESSION OF CHRONIC KIDNEY DISEASE

Once somebody has some degree of kidney disease it frequently progresses over time. The rate of progression can vary. The stages of CKD can be thought of in terms ranging from 1 to 5.

In CKD stage 1, the patient has normal filtering function as measured by the Glomerular Filtration Rate (GFR) of about 120 mls/min. Other levels of CKD are outlined below:

CKD 1 GFR greater than 90mls/min CKD 2 GFR 60 to 89 CKD 3 GFR 30 to 59 CKD 4 GFR 15 to 29 CKD 5 GFR less than 15mls/min

Most patients who are diagnosed as having CKD 1, 2, or 3 have only mild kidney disease and do not progress to ESKD. The eGFR is usually calculated by using the plasma (blood) creatinine level. Once the GFR is below 70, it frequently continues to decline. If the GFR declines by one ml per year, it will take 80 years to progress from CKD Level 1 to CKD Level 5, requiring dialysis or a kidney transplant. However, if GFR declines by 10mls per year it will only take 8 years to go from CKD Level 1 to CKD Level 5.

Please refer to Book 5 for more information.

CAUSES OF KIDNEY DISEASE

here are hundreds of different diseases that can cause chronic kidney disease. Commonly, the condition is due to one of the following:

DIABETES MELLITUS

Diabetes is a disease in which a patient cannot control the amount of glucose in their blood stream. It is caused either by an inability to produce the substance called INSULIN, which controls glucose in the body (Type 1 Diabetes), or, if the body is unable to respond to the insulin that is produced (Type 2 Diabetes). Whether diabetes is treated by insulin, tablets or diet, it can cause kidney disease.

Apart from raised levels of blood sugar, an important feature, of this disease, is the damage that occurs to small blood vessels. The kidneys contain many small blood vessels and, when damaged by high blood sugar levels in the blood, they are replaced with scar tissue and become blocked. Diabetes can also damage the nerves in many parts of the body. When the bladder is affected, it is more difficult to pass urine, resulting in a build-up in pressure on the kidneys, causing further damage.

The urine of people with diabetes has a high sugar content, which encourages the growth of bacteria and, as a result, kidney infections may occur. Poor sugar control combined with high blood pressure can increase your risk of making kidney disease worse.

HIGH BLOOD PRESSURE

Hypertension means high blood pressure.

SYSTOLIC blood pressure is consistently over 140 (systolic is the "top" number of your blood pressure measurement, which represents the pressure generated when the heart beats).

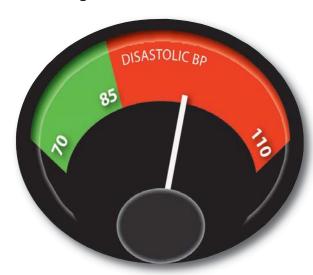
DIASTOLIC blood pressure is consistently over 90. Diastolic is the "bottom" number of your blood pressure measurements, which represents the pressure in the vessels when the heart is at rest). Either or both of these numbers may be too high. Severe high blood pressure can, on its own, cause kidney disease. High blood pressure can make other causes of kidney disease worse.

In any person with high blood pressure, blood vessels (especially small blood vessels) become damaged. Roughly speaking, the higher and the longer blood pressure has been raised, the more blood vessel damage is likely to have occurred.

To control high blood pressure, many patients need medications - known as anti-hypertensives. A lot can be done to control blood pressure including changes in lifestyle, weight loss, exercise, and avoiding salt in the diet.

High blood pressure will commonly require medication to keep blood pressure within the normal range.







NEPHRITIS

The term nephritis covers a group of conditions in which there is long-term inflammation of the kidneys ('neph-' means kidney, and '-itis' means inflammation). Sometimes the condition is described more specifically as glomerulonephritis or GN ('glomerulo-' refers to the glomerulus, which is part of the kidney's filtration unit).

When a glomerulus is damaged, substances not normally filtered out of the blood stream, such as proteins, red blood cells and white blood cells, can pass through the glomerulus and enter the fluid that becomes urine. Progressive damage to the glomeruli can cause urine production to fall and waste products to build-up in the blood, leading to kidney disease. There are many types of glomerulonephritis. These may be grouped as primary and secondary.

In primary GN, only the kidneys are affected. In secondary GN, the kidneys are damaged as part of a more generalised disease that can affect other parts of the body. The exact diagnosis can usually only be diagnosed for certain by a kidney biopsy.

Types of GN include:

FSGS - is scarring within the kidneys that can only be seen clearly under the microscope. Therefore, it is normally only diagnosed after a biopsy of the kidney. The name FSGS comes about in the following way:

- F FOCAL means that some glomeruli are affected but others may not be.
- S SEGMENTAL means affecting only a segment of each glomerulus involved.
- **G GLOMERULO** of the glomeruli.
- S SCLEROSIS meaning scarring.

It seems that the general cause of FSGS is immunological. That is, the antibodies and white blood cells that usually fight off infection cause damage to the body by mistake. This condition commonly results in severe swelling of the legs and high blood pressure. The condition may reoccur after kidney transplantation and cause the failure of a kidney transplant. A number of drug treatments are used to try to stop the damage to the kidney that the condition causes. These drugs include steroids, Cyclosporin, Cyclophosphamide and CellCept.

IgA NEPHROPATHY

This is the most common form of glomerulonephritis in the developed countries of the world. IgA is short for Immunoglobulin A, an antibody which usually helps the body to fight infections and toxins encountered in the gut and lungs. In IgA nephropathy, IgA is deposited in the glomerulus, where they cause inflammation and scarring. In one third of cases, it goes on to cause progressive severe damage to the kidneys. IgA nephropathy tends to be slowly progressive, and so the process of are a type of the kidney disease can take 10 to white blood cell 30 years. present in your

MULTISYSTEM DISEASES
AFFECTING THE KIDNEY

The kidney may be affected in many ways by diseases which are not directly associated with kidney function. This is mainly because the kidneys have a rich blood supply, and come into close contact with all elements of the blood.

SLE (Systemic Lupus Erythematosus)

This is a chronic auto-immune inflammatory disease which affects joints, blood vessels, skin, the nervous system and the kidneys. Inflammation of the glomeruli, within the kidneys, may result in protein and blood loss in the urine and high blood pressure. Kidney problems often occurs within three years of diagnosis of SLE.

AMYLOIDOSIS

Amyloidosis is the term given to a group of chronic disorders characterised by the presence of deposits of an abnormal protein called amyloid. This systemic disease can affect the heart, nervous system, liver and kidneys. Within the kidneys, the amyloid is usually deposited in the walls of the kidney arteries and the glomeruli's blood vessels. This may result in abnormally high levels of protein in urine and can lead to progressive kidney disease.

MULTIPLE MYELOMA

This is a cancer of plasma cells. Plasma cells are a type of white blood cell present in your bone marrow. Multiple myeloma may cause kidney problems, including kidney disease. Higher calcium levels in the blood, due to damage to the bones caused by the myeloma, can interfere with the kidneys' ability to filter your bloods waste. The proteins produced by the myeloma cells can cause similar problems, especially if you become dehydrated.

VASCULITIS

bone

marrow."

This term refers to a group of diseases characterised by inflammation of the blood vessels. Vasculitis causes changes in the walls of the blood vessels,

resulting in thickening, weakening, narrowing and scarring. It is usually due to the body producing antibodies that attack blood vessels throughout the body. Vasculitis may affect blood vessels of any type, size or location and, therefore, may cause dysfunction in any organ system, including the kidneys, lungs, skin and joints. Some of the types of vasculitis which can cause kidney dysfunction include:

- Henoch Schonlein Purpura
- Microscopic Polyangitis
- Polyarteritis Nodosa
- Wegeners Granulomatosis

A blood test called ANCA is commonly positive in vasculitis and is used to monitor the activity of this disease. Vasculitis can be very successfully treated, particularly if diagnosed early and treated with powerful immuno-suppressive medications.

These medications include high doses of steroids, cyclophosphamide, CellCept, and Azathioprine.

POLYCYSTIC KIDNEY DISEASE (PCKD)

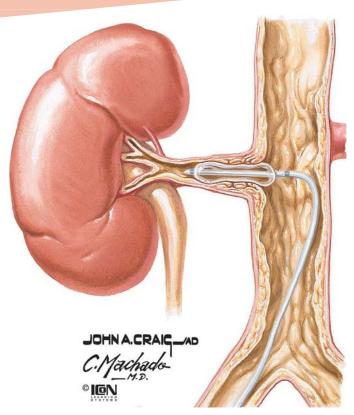
This is a *genetic* disorder characterised by the growth of numerous cysts within the kidney. These cysts are non-cancerous round sacs of water-like fluid. PCKD cysts can slowly replace much of the mass of the kidney, reducing kidney function and leading to kidney disease. People with PCKD may also have similar cysts in their liver, and there is also sometimes an association with a weakness of some of the blood vessels in the brain. PCKD is the most common inherited kidney disease, and children of parents with PCKD have a 50% chance of being affected by it. PCKD frequently causes kidney disease in patients in their 50's. PCKD is diagnosed by a ultrasound scan of the kidneys. Apart from control of blood pressure, there is no "cure" to stop the development of PCKD.

VASCULAR DISEASES OF THE KIDNEY

KIDNEY ARTERY STENOSIS This is a narrowing of the lining of the main artery (blood vessel) that supplies the kidneys. The resulting restriction of blood flow to the kidneys may lead to reduced kidney function and high blood pressure.

This type of hypertension is known as renovascular hypertension and accounts for approximately 5% of patients with hypertension.

Renovascular hyper-tension occurs when the artery to one of the kidneys is narrowed (unilateral stenosis). Kidney disease occurs when the arteries to both kidneys are narrowed (bilateral stenosis). The decreased blood flow to both kidneys increasingly impairs kidney function. This condition may be amenable to treatment with a balloon to open the blocked artery.



OBSTRUCTIVE DISORDERS OF THE KIDNEY

Obstructive Nephropathy - is a kidney disease caused by a blockage to urine flow through the urinary tract. There are many things that can block urine flow. Some of the more common causes include:

- Kidney Stones
- Enlarged Prostate Gland or Prostate Cancer (males)
- Bladder Problems
- Bladder Cancer

Obstruction of the urinary tract results in increased back pressure on the kidneys and increased frequency of urinary tract infections.

These factors cause recurrent episodes of kidney inflammation and scarring, and the kidneys may shrink (atrophy). If the blockage is only for a short time, the kidney can usually recover completely when the blockage is relieved. However, if the blockage is there for a long time (for many days or weeks), it can cause permanent kidney damage.

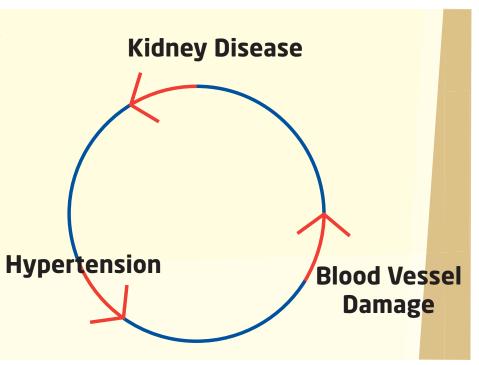
Pyelonephritis This is a bacterial infection of one or both kidneys. Chronic pyelonephritis is kidney damage caused by recurrent or persistent kidney infections. Chronic pyelonephritis is associated with progressive scarring of the kidneys, which can lead to kidney disease. It occurs mainly in patients who have malformations within the urinary tract.

PROGRESSIVE KIDNEY DISEASE

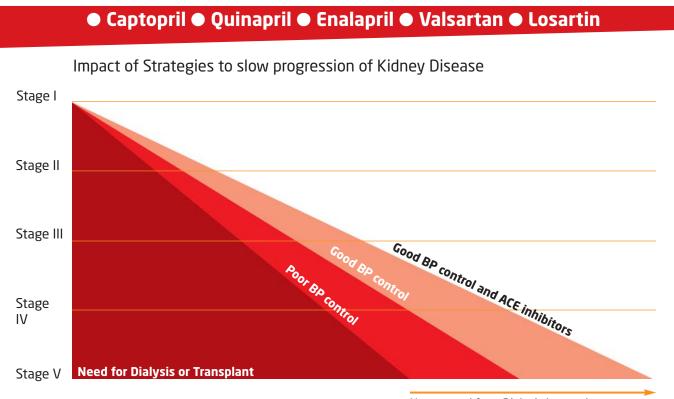
There are many factors that can affect the rate of decline in kidney function.

THESE INCLUDE:

- Underlying cause of kidney disease.
- Blood pressure control
- Use of certain antihypertensive drugs, such as Angiotensin Converting Enzyme (ACE) inhibitors.
- The success of blood glucose control, in patients with diabetes.
- Control of cholesterol



Blood pressure control is one of the factors that can have the biggest influence in slowing the progression of kidney disease. The target blood pressure for any patient with kidney disease should be around 120/80 mmhg. The use of blood pressure medicines called angiotensin, converting enzyme inhibitors or angiotension II blocking inhibitors, has a considerable benefit in slowing progression of kidney disease. Examples of these medicines would include:



SYMPTOMS OF KIDNEY DISEASE

ndependent of the cause of initial kidney injury, kidney function may continue to slowly deteriorate. This is associated with high blood pressure, worsening proteinuria and rising serum creatinine. Many people who have Chronic Kidney Disease (CKD) do not know it because the early signs can be very subtle. It can take many years to go from CKD to End Stage Kidney Disease (ESKD). Some people with CKD live out their lives without ever reaching ESKD.

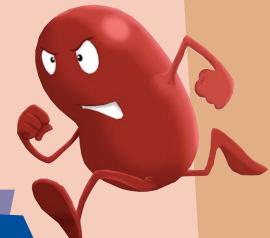
In fact, the majority of people with kidney disease have no symptoms when first diagnosed and are diagnosed on the basis of blood or urine tests.

However, for people at any stage of kidney disease, knowledge is power. Knowing the symptoms of kidney disease can help you get the treatment you need to feel your best.

SYMPTOM 1:

CHANGES IN URINATION

Kidneys make urine, so when the kidneys are failing, the urine may change. How? You may have to get up often at night to urinate. Urine may be foamy or bubbly. You may urinate more often, or in greater amounts than usual, with pale urine. You may urinate less often, or in smaller amounts than usual with dark coloured urine. Your urine may contain blood. You may feel pressure or have difficulty urinating.



WHAT PATIENTS SAY:

"My urine is what I started noticing first.
I would frequently want to go to the bathroom
and when I got there I could only pass a few drops."

SYMPTOM 2:

SWELLING

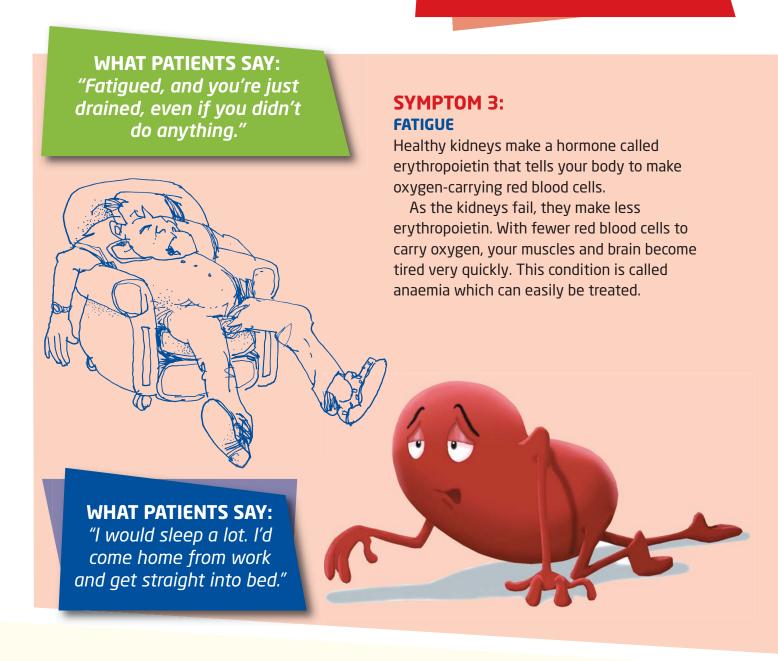
Failing kidneys don't remove enough fluid which then builds up in your body causing swelling in the legs, ankles, feet, face and hands.

WHAT PATIENTS SAY:

"I had a lot of swelling in my ankles. My ankles were so big I couldn't get my shoes on."



Your tests reveal that you are retaining fluids!



SYMPTOM 4:

SKIN RASH/ITCHING

Kidneys remove waste from the blood stream. When the kidneys fail, the build-up of waste products in your blood can cause severe itching.

WHAT PATIENTS SAY:

"It's not really a skin itch or anything, it's just right down to the bone. I was itching and scratching a lot."



WHAT PATIENTS SAY:

"Foul taste in your mouth. Almost like you're drinking iron."

SYMPTOM 5:

METALLIC TASTE IN MOUTH/AMMONIA BREATH

A build-up of waste products in the blood (called uraemia) can make food taste different and cause bad breath. You may also notice that you stop liking to eat meat, or that you are losing weight because you don't feel like eating.



WHAT PATIENTS SAY:

"You don't have the appetite you used to have."

SYMPTOM 6:

NAUSEA AND VOMITING

The severe build-up of wastes in the blood (uraemia) can also cause nausea and vomiting. Loss of appetite can lead to weight loss.

WHAT PATIENTS SAY:

"When I got the nausea, I couldn't eat and I found it difficult to swallow my tablets."

WHAT PATIENTS SAY:

"I couldn't sleep at night because when I lay down flat, I couldn't catch my breath."

SYMPTOM 7:

SHORTNESS OF BREATH

Trouble catching your breath can be related to the kidneys in two ways. Firstly, extra fluid in the body can build up in the lungs. And secondly, anaemia (a shortage of oxygen-carrying red cells) can leave your body oxygen-starved and short of breath.

WHAT PATIENTS SAY:

"You go up a set of stairs and you're out of breath, or you do work and you get tired and you have to stop."



WHAT PATIENTS SAY:

"My memory disimproved a lot I couldn't remember what I did last
week or maybe 2 days ago.
I couldn't really concentrate
on my crossword puzzles
or reading."

SYMPTOM 9:

DIZZINESS AND TROUBLE CONCENTRATING

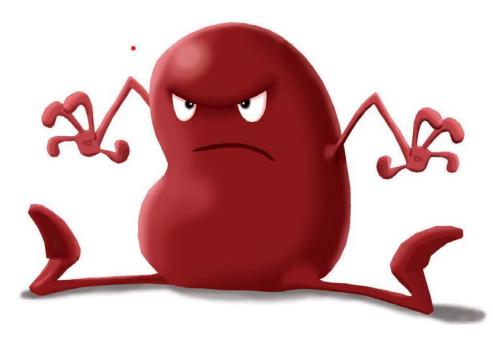
Anaemia related to kidney disease means that your brain is not getting enough oxygen. This, combined with uraemia, can lead to memory problems, trouble with concentrating, and dizziness.

environment.

SYMPTOM 10:

FLANK PAIN

Some people with kidney problems may have pain in the back or side related to the affected kidney. However, most patients with kidney disease do not experience any pain at all.



CHAPTER 3 WHAT ARE DIAGNOSTIC TESTS?

iagnostic tests are used by doctors in order to better assess your health situation. These tests can be used to help establish your initial diagnosis, assess the effectiveness of the treatment you are receiving and/or detect potential complications. Some tests are simple, so simple that often you are unaware that they are happening, like a dipstick analysis of a urine sample. Other diagnostic tests are obvious enough for you to be aware of - like having your blood pressure, temperature and pulse checked. Another group of diagnostic tests considered to be low-risk and non-invasive include x-rays or ultrasound scans.

As soon as the requirements of a specific test involves entry of a needle or a probe into your body, they are considered to be 'invasive'. These tests include procedures such as blood tests, kidney biopsy or x-rays that require an injection of a dye.

WHY ARE DIAGNOSTIC TESTS IMPORTANT?

Diagnostic tests are often an important part of establishing what your health problem is in order that your doctor and nurse can deliver the treatment and care that will best suit your health care needs. The doctor can use other assessment skills to assist in this problem-solving process. These skills involve drawing conclusions from a person's medical history and performing a physical examination. However, a diagnostic test is often required to establish a diagnosis, to plan an intervention, or to monitor progress.

URINE TESTS

What does a dipstick urine test show?

A dipstick urine test, also known as a urinalysis, is a very simple test that is commonly done in clinics or when admitted to the hospital. You will be supplied with a sterile container and asked to carefully pass urine into the container to avoid the sample being contaminated. A nurse will insert, into the urine sample, a special strip that can detect protein, blood, white blood cells and glucose, in the urine, according to the changing colour of the strip.

Protein - this is an important building block in the body. When your kidneys are damaged, protein leaks into your urine (proteinuria). Persistent protein in the urine suggests damage to your kidneys.

Blood - The sample will be examined for colour and clearness. If blood is present, the urine may look red or the colour of cola. This may indicate an infection.

White Blood Cells - Presence in urine indicates infection.

Glucose (sugar) - Presence in urine may indicate diabetes.



WHAT IS URINE CULTURE AND SENSITIVITY?

This is a 'mid-stream' urine sample. Do not collect the first or last part of the urine that you pass, which may contain bacteria or cells normally found on the skin. This is then sent to the laboratory to be examined under a microscope, which can help to diagnose some kidney diseases.

Many things can be seen - like red blood cells. This can be a sign of kidney disease that has damaged the filtering units of the kidney, allowing blood cells to leak into the urine.

It may also indicate other problems such as kidney stones. White blood cells, crystals, and bacteria can also be detected if there are bacteria or white blood cells and this may suggest urine infection. To confirm this, bacteria are allowed to grow on special plates overnight (culture). This allows the doctor to prescribe the most effective treatment for you.



24 HOUR URINE COLLECTION

There are two main reasons to do a 24 hour urine collection. The first is a "urine stone screen" which is used to find the causes of recurrent kidney stones. The second is to measure the eGFR and 24 hour urine protein. However, today we usually use the URPCR instead (See below).

You will be given 1-2 large urine collection bottles and asked to collect your urine over 24 hours. It is important to discard the first urine sample of the day into the toilet and then collect all urine for the next 24 hours into the bottle(s). On the second day, the first sample of urine should be collected. The collection is then complete. A blood test is taken when the urine collection is completed. It is important that the collection is properly carried out or the results will be wrong.

URINE PROTEIN-CREATININE RATIO (URPCR)

Urine protein-creatinine ratio testing is used to detect protein in the urine, to help evaluate and monitor kidney function, and to detect and diagnose early kidney damage and disease. A protein to creatinine ratio is a snapshot of how much protein is present, in the urine, at the time it is collected.

There are many different reasons why there is protein in the urine. This test may be ordered on a random urine sample if a person shows evidence of significant and persistent protein in their urine or has known kidney damage that the doctor wishes to monitor. It is collected by either a 24 hour urine collection or a midstream urine sample.

BLOOD TESTS

Blood tests are regularly carried out to measure how well your kidneys are working.

The table on the next page outlines some of the blood tests that will be performed and what they mean.

BLOOD SAMPLE	NORMAL LEVELS	WHY IT IS BEING MEASURED
Urea (Ur)	2.5-8.5 mmol/L	A waste product produced in the liver and excreted by the kidneys. High values might mean the kidneys are not working as well as they should.
Creatinine (Cr)	49-90umol/L	A waste product produced largely from muscle breakdown. High levels, especially with high Urea levels, indicate problems with the kidneys.
Sodium (Na)	135-145mmol/L	The balance of salt and water in the body.
Potassium (K+)	3.2-5.2mmol/L	Important for proper functioning of nerves and muscles, particularly the heart. High and low levels require medical evaluation. Potassium comes from food, especially fruit, vegetables and nuts.
Calcium (Ca)	2.12-2.62mmol/L	Excreted by the kidney. Important for muscle contraction, cardiac function and blood clotting.
Phosphate (P04)	0.7-1.5mmol/L	Excreted by the kidneys. Necessary for strong bones, teeth, normal functioning of muscle and blood clotting.
Albumin	30-50g/L	Protein in the blood made in the liver. Low levels may indicate that protein is leaking into the kidneys or if someone is malnourished.
RISKS Cholesterol	0.00-5.00mmol/L	Measures how much cholesterol and lipids are present in your blood.
Triglyceride	0.000-1.90mmol/L	
Urate	140-420umol/L	
Parathyroid Hormone	15-65pglm	Concerned with the regulation of extra-cellular calcium levels.
Iron Studies Ferritin	18-240nglml	Main stored iron found in all tissues.
Complete Blood Count (CBC)	13.0-16.0gm/dl	Complete blood count is the red protein in blood. It carries oxygen around the body. One of the kidneys function is to produce erythropoietin (EPO), which stimulates the bone marrow to produce red blood cells. When kidneys fail it may be necessary to give EPO in the form of an injection.
White Cell Count (WCC)	4-11 10^g/l	White blood cells fight infection in the body. Raised/low levels may indicate infection.
Hepatitis B+C	Positive/Negative	Checks for the presence of hepatitis infection.

PLAIN ABDOMINAL X-RAY

A plain abdominal x-ray is used to show the kidneys, urethra and bladder (KUB). It also indicates the size, shape, position and the presence or absence of one or both kidneys.

What preparation is required?

There is no preparation required for this test.



KIDNEY ULTRASOUND SCAN

This is a non-invasive procedure where a transducer (sonar probe) is moved in close contact over the skin over the area of investigation and it can be repeated frequently if necessary. Ultrasound is used to determine the size and shape of the kidneys, to check both kidneys are present, location of kidneys and is useful in detecting cysts.

What preparation is required?

You will be asked to wear a hospital gown. Depending on the area of ultrasound you may be asked to fast prior to the procedure. Your doctor/nurse will inform you if this is necessary.

There are no risks attached to this procedure.

COMPUTERISED TEMOGRAPHY (CT SCAN)

A CT scan, also known as a CAT scan, is a specialised x-ray which provides clear pictures of the inside of your body. In particular, it can give good pictures of soft tissues of the body which do not show on ordinary x-ray pictures.

What preparation is required?

You will be provided with a hospital gown to wear during the procedure. Any jewellery will need to be

removed and depending on what area is being scanned you may need to fast prior to the procedure. Your doctor/nurse can inform you if it is necessary. Also, it may be necessary to drink special fluids prior to the procedure. This is needed if you are having an abdominal/pelvic scan. This drink helps to show the stomach and bowel more clearly. Sometimes a dye (contrast medium) is injected into the bloodstream via a needle. The dye



may give you a flushing feeling and an odd taste in your mouth for a short time. Your doctor will prescribe a course of medication (n-acetylcystine) to take on the day before the scan, the day of the scan, and the day after. This medication protects your kidneys from becoming damaged from the contrast dye that is used during this procedure.

The scan is painless, however, it may take a little time to obtain the necessary pictures.

ECHOCARDIOGRAM

An echocardiogram is an ultrasound scan of your heart. The scan can give clear pictures of the heart muscle, heart chambers and structures within the heart such as the valves. Electrodes will be placed on your chest to allow for a tracing of your heartbeat during the procedure. Some cold gel will be spread on your chest and a transducer is placed on your ribs near your breastbone and directed towards your heart.

The transducer will pick up the echos of the sound waves which transmit them as electrical impulses. The echocardiography machine converts these impulses into moving pictures of the heart. During the procedure you may be asked to breathe in a certain way and change your position.

What preparation is required?

No specific preparation is required and there are no risks attached to having this procedure performed. After the test is completed you can continue as normal.

ANGIOGRAM

An angiogram is a test, using dye and x-ray, to detect if there are any problems in the arteries, valves or chambers of the heart. This test might be performed if you experience tightness or pain in the chest, jaw or arm. A catheter is inserted via the femoral artery (top of the leg) and fed into the bigger artery. The contrast dye is injected and a number of x-rays are taken. You can have an angiogram of your coronary (heart) or renal (kidney) arteries.

Complications of this test include bleeding and a formation of bruising at the catheter site, or occasionally, further deterioration of kidney function.



What preparation is required?

Your doctor will explain the procedure to you and will ask you to sign a consent form to state that you understand the procedure and possible complications. Some complications may include irregular heart rhythms, chest pain, allergic reaction to the dye, bleeding at the groin site and, very rarely, heart attack or stroke.

You will be asked to remove your jewellery and be provided with a hospital gown. It will be necessary to take some blood samples, which will be sent to the laboratory, to ensure it is safe to perform this procedure. Your doctor will also place a needle into your vein (cannula). It may be necessary for you to take medication (n-acetylcystine), prior to this test, to protect your kidneys from the dye that is required for the test.

What will happen after the procedure?

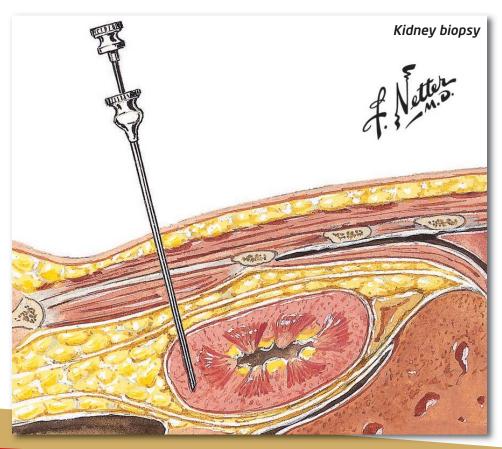
After the procedure, you will be asked to remain on bed rest for 6 hours to allow the puncture site to heal completely. You will need to drink plenty of fluids for the following 24 hours providing you are not on fluid restriction. This will help to flush the dye from your system. A bedpan/urinal will be provided if required. It is very important that you do not bend your leg or sit up before your rest time is over. The nurse will be checking your blood pressure, pulse and the site where the procedure was carried out. Inform the staff if you have pain or discomfort as pain medication can be given.

Your doctor will discuss your results with you before you leave the hospital or at your next appointment. It may be necessary to commence some medication or go for further tests, depending on your results.



KIDNEY BIOPSY

A kidney biopsy is an important test to either establish why the kidneys have stopped working properly or to confirm a diagnosis of rejection in transplant patients.



What preparation is needed?

This procedure is done as a Day Case in the Renal Day Ward. You are admitted to the Renal Day Ward at 7.30am on the morning of your biopsy. There is no need to fast. Your blood pressure will be checked and a blood sample taken to ensure that it is safe to proceed.

The doctors will discuss the complications with you and obtain written consent. The doctor will instruct you on any medications that need to be stopped prior to the procedure e.g. warfarin.

How is the procedure performed?

A kidney biopsy is carried out in the x-ray department or on the ward by the nephrologist. Mild sedation may be given prior to the procedure. If the biopsy is of your native kidney, you will be asked to lie on your tummy.

If the biopsy is of your transplant kidney, you will be asked to lie on your back. This allows the doctor easier access to the kidneys. The skin will be cleaned with antiseptic solution and a local anaesthetic will be given to numb the area. A special biopsy needle is introduced through the skin into the kidney to take the sample. Two or three samples might be required.

What happens after the biopsy?

Following the biopsy, you must stay in bed for 6 hours. This is to allow the puncture site to stop bleeding and help prevent any complications. Following the 6 hour period, you may mobilise gently to the bathroom only. Full mobility can resume the following morning at home if no complications have occurred. Your blood pressure and pulse will be monitored regularly and the biopsy site dressing checked. Each time you pass urine, it should be given to the nurse, who will then test it to see if there is any bleeding. Painkillers will be prescribed if you need them.

You will be able to eat and will be encouraged to drink plenty of fluids (providing you are not on fluid restriction). Usually, you will be discharged the same day.

"Following
the biopsy, you
must stay in bed for
6 hours. This is to allow
the puncture site to
stop bleeding and help
prevent any
complications."

It is advisable to avoid any strenuous exercise for a week after the biopsy to reduce the risk of bleeding. After this there should be no reason why normal activities cannot be resumed.

If you experience severe pain over the biopsy area or notice blood in your urine, you should report back to your doctor.

What are the possible complications during a kidney biopsy?

Any medical or surgical procedure carries risks. Patients are asked to undergo procedures because it is felt that the benefits outweigh the risks. Complications of kidney biopsy are rare. The most important is bleeding, and you are closely monitored after the biopsy to detect bleeding.

You may have pain or discomfort after the biopsy. Painkillers can be taken to reduce any discomfort. If you experience severe pain, after the biopsy, you should contact the kidney unit.

Other possible complications of kidney biopsy include:

- Bleeding into the kidney, which can result in loss of the kidney;
- Persistent haematuria (blood in your urine);
- Biopsy of an organ other than the kidney;
- Rupture of the kidney;
- Death (**extremely** rare).

When do I get the results?

It takes up to 48 working hours for the laboratory to give a preliminary report and about a week to ten days to get a full written report. Your doctor will discuss the results of the biopsy with you and discuss appropriate treatments if required.

CHAPTER 4

ietary treatment is an important aspect of care for all patients with kidney disease. It is necessary to meet with your dietician to discuss individual needs for your renal diet.

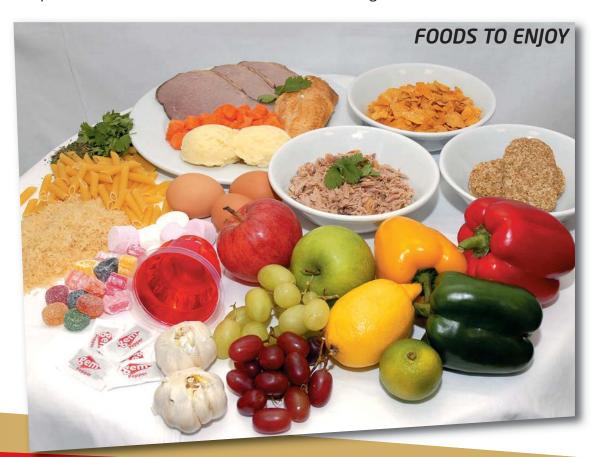
The following information will give you an overview of a renal diet, prior to dialysis, and also what to expect when you commence dialysis. You will be informed by the dietician when it is necessary to commence this diet. All patients have different dietary needs, so it is important to start this diet only under the guidance of the dietician.

A healthy balanced diet contains correct amounts of protein, carbohydrate, fat, vitamins and minerals. It is important that your diet is balanced and varied to keep you in optimum health.

Some of the main functions of the kidney that relate to the diet include:

- Excretion of waste products;
- Control of fluid volume in the body;
- Control of blood pressure.

When food and drinks are consumed, our bodies use what is needed and the rest is turned into waste products which can be excreted as urine. When your kidneys are not working properly, these waste products can build-up in your blood and cause complications, which will be discussed in the following sections.



SALT

Salt is an important aspect of dietary treatment at all stages of your kidney disease. High intake of salt, from the diet, can cause problems with blood pressure control and fluid retention. It is advised to avoid adding any salt to meals and also to reduce the intake of very salty foods such as processed meats, bacon, sausages, soup and packet sauces. Your dietician will advise vou on suitable alternatives to using salt.



PROTEIN

Protein intake from the diet is important during the progression of chronic kidney disease and also when you commence dialysis. The protein we eat is used for tissue repair and growth. Any unused protein is broken down into waste products, including urea and creatinine. As your kidneys are unable to excrete urea and creatinine properly, they build up in your blood and cause symptoms such as nausea and loss of appetite.

By eating large amounts of protein foods e.g. meat, fish, chicken, eggs, cheese, milk and yoghurt before commencing dialysis, you will affect the build-up of urea and creatinine in your blood. An appropriate daily intake of protein should be advised by your dietician.

However, once dialysis treatment has commenced it is important to make sure that your body is getting enough protein to prevent malnutrition. Some of your stores of protein are lost during the Haemodialysis or Peritoneal dialysis sessions.

How much protein you need depends on your body size and is specific to each individual.

PHOSPHATE

Phosphate is another mineral found in many foods, mainly meat and dairy products such as milk, cheese, yoghurts, and also bran nuts and cola. Calcium and phosphate work together to keep your bones, teeth and blood vessels healthy. When phosphate and calcium levels are elevated, or out of balance in kidney disease, the extra calcium and phosphate join together to form hard deposits in your body. This is known as calcification. These deposits can form in the heart, lungs, blood vessels, joints and other soft tissue. High phosphate levels also affect your bones, causing kidney bone disease. Over time bones become brittle, weak and painful and liable to fracture easily.

As with potassium, an elevated phosphate level will require you to reduce the intake of phosphate from your diet. It may also be necessary to take phosphate binding substances with your food to reduce the absorption of phosphate from the gut.

POTASSIUM

Potassium is a mineral found in many foods, mainly fruits, vegetables and potatoes. It is necessary for muscle contractions but a very high level in the blood can be dangerous as it may cause irregular heart rythym. If your potassium levels increase, above normal, you will need to avoid certain foods that are high in potassium as advised by your dietician. High sources of potassium in the diet include bananas, dried fruit, peas, beans, spinach and potato products such as chips and crisps.

The dialysis diet provides enough potassium to meet the needs of your body, while preventing accumulation between dialysis sessions.

FLUIDS

If you are treated with haemodialysis or peritoneal dialysis you may need to limit your fluid intake. The amount of fluid you will be allowed depends on the amount of urine you produce. You can reduce your fluid intake by using small cups/ glasses, and spacing out your drinks throughout the day. You can also help prevent thirst by limiting the amount of salt and salty foods that you eat.

Each person, with kidney disease, is very different and so are their needs and requirements. The dietary advice you are given depends on a number of factors including the stage of kidney disease, the type of treatment you are on, your blood results, your body weight, and the presence of other medical conditions e.g. diabetes mellitus, high cholesterol levels. The dietician will, therefore, provide you with information that is designed for you as an individual to suit your own specific needs.

USEFUL RESOURCES





Over 100 special recipes created by Ireland's top chefs, for adults living with kidney disease. Perfect for entertaining or for planning that special meal that is suitable for all to enjoy.

ORDER YOUR COPIES NOW.

www.ika.ie

3

www.irishkidneydiet.ie

A new website by Renal Dietitians to help you discover what you can eat when you need to follow a kidney diet









Information
Recipes
Meal plans
Shopping lists
Dietitians Blog
Videos
FAQ's



This website has food, diet and nutrition information for people living with chronic kidney disease (CKD), and their carers. Discover what you **can** eat when you need to follow a kidney (renal) diet.

GUIDE TO SUITABLE FOODS TO USE WITHIN YOUR RENAL DIET:

Foods Low in Potassium	Foods High in Potassium
✓ Apples, pears, plums, mandarins, grapes	x Banana, dried fruit, prunes, apricot,
✓ Kiwi, peach	* Rhubarb
✓ Cauliflower, peppers, carrots, broccoli	★ Peas, beans, mushrooms, spinach, beetroot
✓ Cabbage, green beans, turnip	≭ Salt substitutes e.g. Losalt.
✓ Boiled/mashed potato, rice, pasta	★ Jacket/chipped/roast potato
✓ Spirits, white wine, boiled sweets	* Beer, stout, red wine, chocolate, coffee

Foods Lower in Phosphate	Foods High in Phosphate
Rice milk, ask your dietitian for more information	Milk (including nut milks), yoghurt, cheese
Porridge, Weetabix, Cornflakes, tea	Bran cereals, peas, beans, corn, nuts
Marshmallows, jellies, boiled sweets	Toffee, cola, chocolate

Foods to ENJOY	Foods to AVOID
Pepper, herbs, garlic, mustard, lemon	Salt, stock cubes, soup
Home-made stock	Salted snacks
Chicken, lamb, pork, beef, eggs	Processed meats
Fish (excluding smoked fish)	
BREAKFAST	
Boiled/poached/fried eggs	Bacon, sausage, black & white pudding
Plain omelette, bagel, English muffin	Salt substitutes
Croissant, french toast, porridge,	Bran cereals, muesli
Cereals (excluding bran cereal & muesli)	
SANDWICH IDEAS	
Turkey and cranberry sauce	Ham, corned beef, salami
Roast beef & mustard	Cheese
Egg mayonnaise, chicken, tinned salmon (no bo	ones)
MAIN COURSE	
Ask for sauces/gravy to be served	Casseroles, cured or salted meats
on the side and use sparingly	
Roast/grilled pork/lamb/beef	
Chicken/turkey or fish	
SIDE ORDERS	
Green beans, cabbage, asparagus	Spinach, mushrooms, peas, corn
Carrots, cauliflower, broccoli	Chipped/jacket potato
Plain rice, pasta, noodles, couscous	
DESSERT	
Canned peaches, pears, fruit cocktail	Dried fruit, fresh fruit cocktail, melon
Fresh grapes, fresh & canned pineapple	Banana, orange
Jelly, plain and cream cakes, apple tart	Desserts with chocolate, nuts, dried fruit
Sherbet, sorbet, plain biscuits, Pavlova	Coconut, milk pudding, ice cream

CHAPTER 5



MEDICATIONS

eople with Chronic Kidney Disease (CKD) will find themselves on a variety of medications. These medications can be classified according to what they do and include the following:

These are just examples of some of the drugs available. Medications have different brand names so it is important to realise that different coloured medication may be given while in hospital.

- Phosphate binders
- Antihypertensive medications
- Diuretics
- Erythropoietin (EPO)
- Oral and intravenous iron

PHOSPHATE MEDICATIONS

Many of the foods you eat contain an element called phosphate. Foods particularly high in phosphate include all dairy products such as milk and also chocolate, pizza and bread. When kidneys fail they are not able to get rid of phosphate and it will build up in your body. It is necessary to take medications called 'phosphate binders'. There are a number of different phosphate binders available. They all work in the same way in that they bind phosphate in your stomach, which prevents the phosphate being absorbed into your circulation. **All phosphate binders are taken with meals.** This is the only way these tablets work and are of no benefit at any other time.

The available phosphate binders include:

Calcium containing compounds. These calcium tablets are effective at binding phosphate but, sometimes, can cause the calcium level in the blood to go too high. These tablets may cause constipation or gas. They are chewed and taken with meals.

Sevelamer (Renagel). This tablet is swallowed and not chewed. Frequently you may have to take as many as 3 tablets, 3 times a day. This tablet may also cause some constipation or abdominal discomfort. It has the advantage over other phosphate binders in that it does not contain calcium.

Foznol is also another phosphate binder commonly used.

BLOOD PRESSURE MEDICATIONS

It is very important to control blood pressure - aiming to keep blood pressure as close to normal as possible. There are many approaches to control blood pressure, including weight loss, salt avoidance, exercise and reduction in alcohol intake. Many patients will, however, also need to take blood



pressure tablets. There are many different classes of blood pressure tablets including:

- Betablockers
- Calcium channel blockers
- Angiotensin converting enzyme inhibitors (ACE)
- Angiotensin receptor blockers (ARB)
- Alpha blockers

BETABLOCKERS

(e.g. Atenolol, Metoprolol, Bisoprolol)

Betablockers are amongst the most commonly prescribed blood pressure medications. Examples include Atenolol and Metoprolol. These tablets are used to treat cardiac conditions in addition to high blood pressure. Betablockers work by blocking the effect of adrenaline on blood vessels and the heart. Consequently, they will slow the heart a little. People with severe asthma need to be careful with these medications as they may make asthma worse and they may also, occasionally, make your hands feel cold. Occasionally, betablockers may give you nightmares.

CALCIUM CHANNEL BLOCKERS

(e.g. Istin, Dilzem)

Calcium channel blockers work by directly making blood vessels relax by blocking calcium going into muscles around the blood vessels. The commonest sideeffect of these medications is swelling of the legs. They may also, occasionally, cause redness of the skin.

ALPHA BLOCKERS

Alpha blockers work by blocking the function of adrenaline on blood vessels. Occasionally these medications may cause you to become dizzy when you stand up.

ANGIOTENSIN CONVERTING ENZYME INHIBITORS (ACE) AND ANGIOTENSIN RECEPTOR BLOCKERS (ARB)

One of the important hormones in the body that controls blood pressure is called angiotensin. ACE inhibitors work by blocking the production of this hormone and ARB work by stopping the effect of angiotensin. These drugs are used to treat blood pressure and heart failure. They are also effective at slowing the decline in kidney function. ACE inhibitors may sometimes cause a persistent cough which resolves when the ACEinhibitor is stopped. These medications may also cause the potassium levels to rise too high. Your bloods will be carefully monitored for this.

DIURETICS

(e.g. Lasix, Burinex)
Diuretics are
medications that
make you pass
more urine. These
medications are
used to treat
swelling of the legs,
heart failure and
are also commonly



used to treat high blood pressure. Diuretics can also be prescribed for patients who are on dialysis to help produce more urine and gain less weight between dialysis treatments.

These medications will work for up to six hours after they are taken. Side-effects of diuretics include dehydration if they make you pass too much urine.

VITAMINS

When you have kidney disease you will have lower levels of vitamins in your body than you need, so you may need to take a number of different vitamins including:

- Multivitamins:
- Folic acid;
- Vitamin D.

MULTIVITAMINS

Patients, who are on dialysis, commonly lose much of the water soluble vitamins through the dialysis machine. It is important that you do not take just any multivitamins as some of these preparations may contain too much of the fat soluble vitamins.

Folic acid is important for many of the metabolic functions of the body.

VITAMIN D

The kidney is responsible for converting the vitamin D that is in your body into the active form of vitamin D. People with kidney disease lack the ability to produce the active form of vitamin D. Vitamin D is important for the absorption of calcium and for bone health which is important to prevent

glands the parathyroid becoming overactive. mentioned earlier the parathyroid (PTH) glands are small glands in your neck that control calcium levels in your body. People with kidney disease can have PTH levels that are too high. One of the major functions of Vitamin D is to lower PTH levels. Too much vitamin D may cause the calcium, in your blood, to go too high so your doctors and nurses will carefully monitoring calcium levels by obtaining regular blood samples.



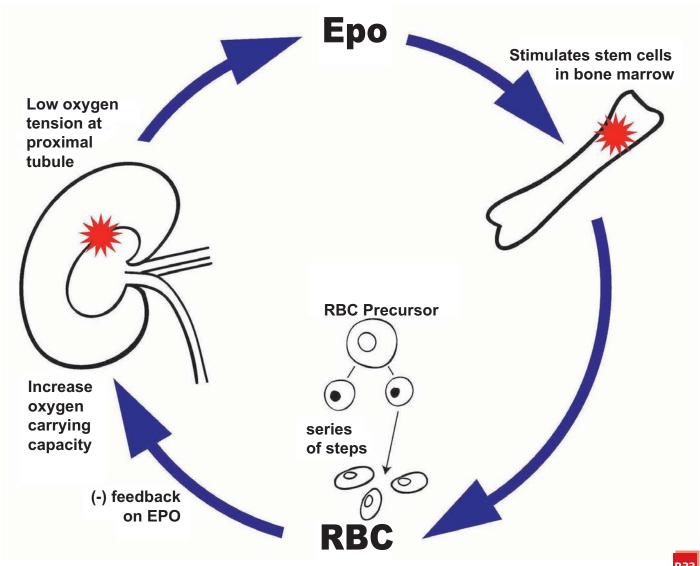
ERYTHROPOEITIN (EPO)

Patients with kidney disease very often have a low red cell blood count. Under normal circumstances kidney produces a substance called the erythropoietin (EPO). EPO works on the bone marrow cells to make more blood cells. EPO needs to be administered by way of an injection. It cannot be taken as a tablet. The injection may be given as an injection directly into the dialysis machine or as an injection into the skin. EPO can be administered either once, twice or three times a week or sometimes every second week or, even only once a month, depending on your blood. Patients will feel much better on EPO as they will acquire more energy with a higher haemoglobin.

EPO is prescribed on a high-tech prescription.

Your chemist will give you 4 or 5 vials, which you will need to keep in the fridge (not the freezer). Your kidney nurse will either teach you how to administer the medication yourself or arrange a district nurse or your GP practice nurse to administer it. It is also possible to have your injection when you attend for dialysis.

It is important to stay on this medication until your doctor indicates otherwise. You will need frequent blood tests to monitor the effectiveness of the EPO as too much EPO may cause your blood to go too high or you may need to increase the dose if you are not getting enough benefit. It is important that you bring your medication with you on admission to hospital or for each dialysis treatment if required.



ORAL OR INTRAVENOUS IRON

EPO is extremely effective at increasing your blood count. However, in order for the EPO to work effectively your body needs to have enough iron. It is difficult for your body to take in enough iron through regular diet for the EPO to reach full effect. Your doctor may prescribe oral iron. Iron tablets can be a little unpleasant to take as they may make your stomach upset or cause you constipation. Iron tablets will also

make your bowel movement black and tarry.

If you cannot tolerate iron tablets or your doctor determines, by blood test, that you are still low in iron, you may be prescribed intravenous iron. If you are on haemodialysis, you will be given iron directly into the dialysis circuit.

If you are on home dialysis or not on dialysis, you will need to come to Renal Day Care in the Renal Centre, for infusions of iron treatment.



Intravenous iron is a safe drug but there is an occasional occurrence of an allergic reaction. So, before you get the first dose, you will generally receive a test dose (small dose).

PARATHYROID DRUGS

As we discussed under Vitamin D, your parathyroid glands are four small glands in your neck that control the level of calcium in your blood. In kidney disease, parathyroid (PTH) hormone tends to increase. When this happens, it can have a number of effects, including weakening of your bones



and causing pain in your bones and joints. In the first instance, your doctor may give you vitamin tablets (Rocaltrol or One Alpha); however, if these do not work, it may be necessary to receive Vitamin D intravenously (Zemplar) directly into the dialysis machine. If these strategies are not adequate, your doctor may advise you to take another drug called Cinacalcit (Mimpara). This drug is extremely effective at lowering PTH level. The main side effect of Cinacalcit is that it may cause a little nausea after it is taken. This, generally, settles down after a few days.

Occasionally, if these drugs are not effective in controlling your PTH level, your doctor may recommend an operation to remove part of your parathyroid glands (a parathyroidectomy).

CAUTION WITH COMMONLY USED DRUGS

When you have kidney disease, you need to be very with careful all vour medications. You need to make sure your doctor knows that you have kidney disease before he prescribes you any medications or alters your dose. Only take medication that is prescribed for you. Please inform your doctor or chemist if you plan to take herbal medication as some of these may interact with your regular medication. Similarly, you need to be very cautious of many 'over-the-counter' medications

which are easily available. Some of these include:

 NSAIDS or anti-inflammatories should be avoided. Examples of these include: Ponstan, Neurofen, Ibuprofen, Advil, Difene, Voltarol, etc.



For pain control you should take simple analgesics such as Panadol or Solpadeine. If you need something stronger, speak with your doctor.

It is very important that you know your medications and how they work. Your healthcare team will work alongside you to achieve this.



DRUGS TO AVOID

- **X** Ponstan
- **X** Neurofen
- **X** Ibuprofen
- **X** Advil
- **X** Difene
- × Voltarol

CHAPTER 6



CAUSES OF INFECTIONS

The common causes of Infections in Ireland are:

- Bacteria (e.g., salmonella, TB, E.Coli).
- Viruses (e.g., common cold, flu, winter vomiting bug).
- Fungi (e.g., thrush, athletes foot).

Infections are spread by:

- Air (e.g. TB, chicken pox).
- Droplet, sneezing and coughing (e.g., mumps, rubella, common cold).
- Direct contact (e.g., salmonella from eating uncooked chicken, and sexually transmitted diseases such as syphilis).
- Indirect contact (e.g., salmonella from a sandwich made by the unwashed hands of a person infected with salmonella).
- Vectors e.g., mosquitoes spreading malaria.

PREVENTING INFECTION

The human body has developed general and specific defences against infection. General defences protect the body against all infections and examples include skin, secretions such as tears, cilia (tiny hairs), which filter air entering the lungs, and body washings, such as flow of urine from the bladder, which washes away bacteria with the urine.

Specific defences develop when the body's immune system produces antibodies against certain diseases. These antibodies develop after an infection (e.g., chicken pox) or after vaccination (e.g., whooping cough, rubella) and ensure that infection or re-infection very rarely occur.

People with kidney disease have an immune system that does not work as efficiently as normal. In addition, some complications of common illnesses such as pneumonia after flu can be dangerous for people with kidney disease. However, everyone can assist his or her natural immune system preventing infection by:

- Eating a well balanced diet and taking regular exercise
- Good general hygiene will help keep the skin in good condition
- Regular hand washing, especially before eating, and after using the toilet
- Attending for regular check-ups with your kidney specialist and GP will ensure that your kidney function and general health are maintained
- Attending your GP or the kidney unit promptly if you are not feeling well

INFECTIONS AND KIDNEY DISEASE

People with kidney disease are vulnerable to the same infections as the general population, such as flu, measles and mumps. However, they are more vulnerable to certain infections due to the treatments used (haemodialysis, peritoneal dialysis and transplantation) and due to regular hospital admissions (MRSA, VRE, and C.Difficle).

Infection complications associated with haemodialysis treatment

Haemodialysis treatment is known to be a risk for:

- 1. Bacterial infections associated with access, i.e., catheters, fistulas and grafts.
- 2. Blood borne viral infections (hepatitis B, C and very rarely HIV).

Access infections

See chapter 4 (Book 2 - Haemodialysis and Peritoneal Dialysis - A Guide for Patients) for detailed information on infections associated with access.

Blood borne infections associated with haemodialysis (HD)

Outbreaks of viral blood infections (Hepatitis B and C) have happened in haemodialysis units. As a result, our unit takes infection control very seriously indeed and makes every effort to keep the risk to an absolute minimum.

The measures include the following:

- All patients are screened on admission and routinely for hepatitis B & C & HIV
- All staff are vaccinated against hepatitis B
- All patients are strongly recommended to be vaccinated against hepatitis B
- Patients with known infections are treated in single rooms, on special machines.

All equipment used, on each patient, is either disposed of after each use or cleaned and disinfected after every use.

Infection complications associated with Transplantation

See chapter 8 (Book 3 - *Kidney Transplantation* - *A Guide for Patients*) for information on infections, associated with transplantation.

MRSA, VRE, CRE/CPE AND C.DIFFICLE

People who have regular hospital admissions, such as kidney patients, are at increased risk of acquiring MRSA, VRE, CRE/CPE and C.DIFFICLE.

MRSA

What is MRSA?

MRSA is the shortened term used when referring to **M**eticillin **R**esistant **S**taphylococcus **A**ureus. Staphylococcus aureus (S. aureus) is the name of the bacteria. The sensitive strain is found in the nose and on the skin of 20-30% of healthy people. The resistant strain (MRSA) means that it cannot be treated with antibiotics normally used to treat the sensitive strain.

Where is MRSA found?

MRSA is most often found in hospitals or nursing homes where antibiotics are used frequently therefore encouraging the development of resistant strains of bacteria.

How does a person acquire it?

MRSA is transferred from one person to another by human contact. The main method is on hands, during patient care. Patients, who are carriers, may pass it on to other patients if they are in close contact.

Does MRSA make a patient more ill?

Some patients are colonised with MRSA and others have infections caused by MRSA. A patient is colonised with MRSA when he/she has no signs or symptoms of infection. It does not alter their treatment and is not a reason to stay in hospital.

MRSA infection, like other infections, varies from mild to severe and depends on other factors, such as where the infection is and the patient age and underlying conditions. A person found to be colonised or infected with MRSA will be nursed separately from other patients, in a single room (isolation), or in a room with others who also have MRSA (co-horted).

What is the treatment for MRSA?

A patient, colonised with MRSA, is treated with special washes and ointments.

A patient, infected with MRSA, is treated with antibiotics, in tablet or by a drip into a vein.

VRE

What is VRE?

VRE is the short-term used when referring to Vancomycin Resistant Enterococci. Enterococci are bacteria found in the faeces of humans. Most of the time enterococci are part of the normal bacteria of the bowel and do not cause disease. A strain of enterococci has developed resistance to vancomycin, which is an antibiotic used to treat serious infections including MRSA infections.

Where is VRE found?

VRE is found in hospitals where patients are very unwell, such as intensive care, kidney and transplantation wards. Enterococci can survive on surfaces, ledges and floors.

How does a person acquire VRE?

VRE may be transferred from one person to another by direct contact, particularly from hands, during patient care.

Does VRE make a person more ill?

This varies from patient to patient. The majority of patients are colonised, whilst some are infected. Colonised means that the VRE is not causing infection. The presence of VRE colonisation does not alter their treatment and is not a reason to stay in hospital. VRE infections can vary from mild to severe and depends on factors such as the site of the infection and the patient's overall condition.

Patients with VRE in a wound or in a urine specimen or those having diarrhoea need to be nursed in a single room (isolation) or nursed in a room with other patients with VRE (co-horted).

What is the treatment for VRE?

Infection, with VRE, is treated with antibiotics usually given via a drip into a vein. Colonisation with VRE does not require any special treatment.

CRE (Carbapenem Resistant Enterobacteriaceae) ALSO KNOWN AS CPE (Carbapenemase Producing Enterobacteriaceae)

What is CRE/CPE?

CRE/CPE are bugs (bacteria) that live in the bowel and that cannot be treated by certain antibiotics, known as carbapenem antibiotics. In most people, CRE/CPE bugs are harmless and do not cause infection. However, if a patient is prone to infection and the infection is caused by CRE/CPE, it can be difficult to treat because many of the commonly used antibiotics will not work against CRE/CPE.

Can CRE/CPE be harmful?

For most patients, CRE/CPE lives harmlessly in the bowel and does not cause infection (this is called 'colonisation'). This is because the person's immune system keeps CRE/CPE in check in the bowel and prevents it from spreading elsewhere in the body. Sometimes however, CRE/CPE can cause infection in patients, for example when they need intensive care or while receiving chemotherapy. CRE/CPE can cause infections, such as kidney infections, wound infections or in severe cases, blood infection. Antibiotics are needed to treat CRE/CPE infection. Doctors need to know as soon as possible if a patient is carrying CRE/CPE so that the most effective antibiotics can be chosen to treat the infection.

How do people get CRE/CPE?

 Patients who have already taken lots of antibiotics are more at risk of picking up CRE/CPE. The reason for this is that the more bugs are exposed to antibiotics, the more likely they will develop 'resistance' to that antibiotic so that antibiotic no longer works.

- CRE/CPE is more common in certain countries than others - if you have been a patient in a hospital abroad in the past year, you need to let your doctor know, especially if you are being admitted to an Irish hospital so that they can test for CRE/CPE.
- CRE/CPE is carried by patients, healthcare staff or visitors, either harmlessly or if they are infected with it. It can spread between patients through direct contact with each other or by touching items or surfaces that the person with CRE/CPE may have touched such as bed rails, toilets or equipment. As patients in hospital are much more vulnerable to infection than patients in their own homes, special precautions are required to prevent the spread of CRE/CPE between patients in hospital.

What are the special precautions for patients with CRE/CPE?

The precautions are designed to prevent CRE/CPE spreading between patients on the ward. Anybody found to have CRE/CPE is placed in an isolation room with their own toilet or commode. Staff or visitors must wear gloves and aprons or gown before entering the isolation room and before coming into contact with the patient to protect their hands and clothes from CRE/CPE. Patients, staff and visitors must pay special attention to hand hygiene. All staff must clean their hands before and after any contact with every patient, regardless of whether the patient has CRE/CPE or not.

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Why do we need to know if a patient is carrying CRE/CPE?

It is very useful to know if a patient is carrying CRE/CPE before it makes them sick. If it is known beforehand that the patient is carrying CRE/CPE then the most effective antibiotics can be chosen early to treat any signs of an infection. When staff know that a patient is carrying CRE/CPE, special precautions can be put in place immediately, to stop the CRE/CPE spreading to other patients, some of whom might be particularly prone to developing infection if they came into contact with CRE/CPE.

Who should be tested for CRE/CPE?

- CRE/CPE is a very common bug in some countries. If you have been admitted to hospital in another country in the past 12 months, you should tell your doctor or nurse so that a CRE/CPE test can be done.
- Some Irish hospitals have also picked up CRE/CPE in patients. If you have been admitted to an Irish hospital in the past 12 months, you should tell your doctor or nurse. They will then be able to decide if you need to be tested for CRE/CPE. Your nurse or doctor will know which Irish hospitals have reported CRE/CPE cases.
- Some hospitals may want to perform a CRE/CPE test on patients who might be prone to developing infections, for example those needing intensive care treatment.

How do you know if a patient is carrying CRE/CPE?

A swab from a patient's back passage is the quickest and easiest way to check for CRE/CPE, as it is usually detected in the bowel. Your nurse will take the swab and send it to the laboratory to be checked for CRE/CPE.

How long does it take for CRE/CPE results to come back from the laboratory?

The laboratory will usually be able to rule out CRE/CPE within three days. However, if CRE/CPE is suspected, the sample will be sent to a specialist laboratory for more detailed testing. These results

may take one or two weeks and you will be given your results as soon as they are ready. You will stay in an isolation room until the final results are received. If you don't have CRE/CPE, you may be moved out of the room, unless there is another reason for staying in isolation. If CRE/CPE is picked up in the sample, you will stay in the isolation room for the duration of your hospital stay, to prevent spreading CRE/CPE to other patients.

What happens when I'm ready to leave hospital?

You can go home as soon as your doctor says you are ready, even if you have CRE/CPE. Carrying CRE/CPE will not affect your discharge.

How can I help?

If you go to the doctor, dentist, another clinic, hospital or nursing home, let them know you have had a positive CRE/CPE result. They can then take special precautions to stop CRE/CPE spreading to other patients who might be more prone to getting a CRE/CPE infection.

What happens when I go home?

There is no need to take special precautions at home. People in the community are usually fit and healthy so they are less likely than hospital patients to pick up CRE/CPE. Clothes, bed linen and dishes can be washed as usual. It is always very important to wash your hands carefully after using the toilet and before preparing meals or eating to stop bugs spreading to other people. Clean hands protect you and others from lots of infections, not just CRE/CPE.

Will I get rid of CRE/CPE?

Your body may clear CRE/CPE from the bowel as you recover, but this is not always the case, and it may remain in the bowel for some time. Taking antibiotics can encourage CRE/CPE to grow in the bowel again, so they should only be taken for a very good reason. Antibiotics are available to treat CRE/CPE infection but they will not clear CRE/CPE from the bowel.

C. DIFFICLE (CLOSTRIDIUM DIFFICILE) What is Clostridium Difficle (C. Diff)?

C. Diff. is a bacteria that causes diarrhoea and may cause intestinal conditions such as colitis. It is a common infection in hospitals and long-term facilities.

The use of antibiotics alters the normal bacterial content of the bowel and, thereby, increases the risk of developing C. Diff. diarrhoea.

Where is C. Diff found?

C. Diff is found in the bowel of some people and can also survive for a long time on surfaces.

How do people get C. Diff?

Healthy people are not at risk from getting C. Diff. People who have other illnesses or

conditions requiring prolonged use of antibiotics and the elderly are at risk of infection. They can become infected if they touch items that are contaminated and then touch their mouth.

Does C. Diff make a person more ill?

In most patients, the symptoms are mild and discontinuing treatment with antibiotics and fluid replacement results in rapid improvement. Sometimes, it is necessary to give a specific antibiotic, by mouth, for the condition. Unfortunately, 20-30% of patients relapse and need further courses of antibiotics.

Patients need to be nursed in a single room (isolation) or, in a room with other patients with C. Diff (cohorted), until bowel movement has returned to normal.

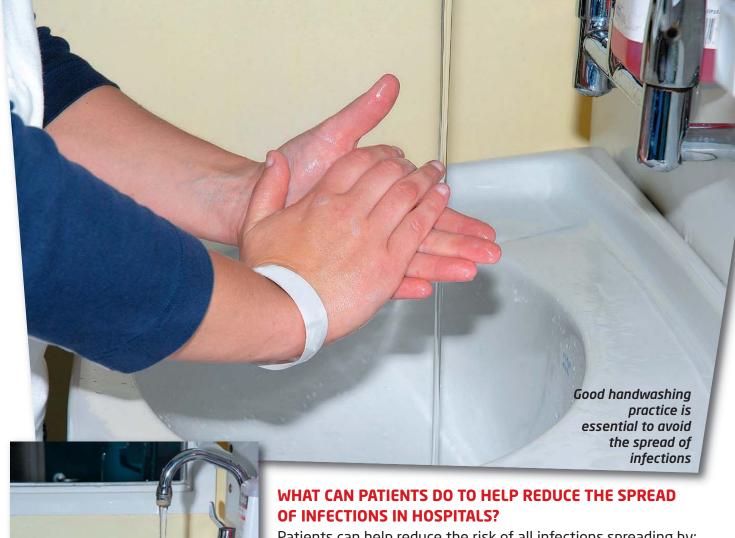
VACCINATIONS RECOMMENDED FOR PEOPLE WITH CHRONIC KIDNEY DISEASE
As prevention is always better than cure, the Department of Heath and Children
advise that certain vaccinations be given to people with kidney disease. Your kidney
doctor or GP will advise you when you need to start getting vaccinated, but, in
general, once a diagnosis of chronic kidney disease is confirmed, the vaccinations
listed below should be given:

- Pneumococcus This bacterium can cause serious infection in the lungs (pneumonia), the blood (bacteraemia) and covering of the brain (meningitis). Vaccination consists of a single injection, followed by a once-off booster dose 5 years later.
- Influenza (flu) An annual flu vaccine is advised, as infection can be complicated by pneumonia, which is dangerous for people with chronic illness.
- Hepatitis B Hepatitis B is a serious illness and as haemodialysis is a recognised risk for acquiring Hepatitis B, vaccination is advised. The vaccination course varies, depending on the product used, but it is usually 3 or more injections, over a 6-month period, with a follow-up blood test to check if immunity has developed. Some people need an additional

- injection (boost) or a repeat course to develop immunity. In addition, people on haemodialysis or peritoneal dialysis have a blood test, yearly, and, depending on the result, may need a boost.
- Varicella (chicken pox) Vaccine for patients not immune and planning to receive a transplant.

People should not get the vaccines if they ever had a life-threatening allergic reaction to yeast (Hepatitis B), eggs (flu) and/or to a previous dose (all vaccines). Pregnant women should discuss vaccination, with their doctor, and people who are ill should defer vaccination until feeling better.

While a vaccine, like all medicines, is capable of causing a serious problem, such as severe allergic reaction, the risk of vaccinations causing serious harm, or death is extremely small. **Getting vaccinations is much safer than getting the disease.**



Patients can help reduce the risk of all infections spreading by:

- Washing hands or using alcohol gel after using the toilet and before meals.
- Reminding staff to wash their hands, or use alcohol gel before they care for you.
- Advising visitors who are feeling unwell not to visit.
- Advising visitors to wash their hands before and after visiting and to avoid going from one ward to another during visiting
- Seeking advice from ward staff if young children wish to visit.
- Complaining to the ward sister/consultant or any staff member if the general ward hygiene is not satisfactory or if staff are not washing their hands.

PREVENTING THE SPREAD OF ALL INFECTIONS IN HOSPITALS

This hospital, along with all hospitals in the country, is working hard to reduce the spread of all infections in hospitals by:

- Improving hygiene throughout the hospital;
- Improving hand hygiene of staff and patients;
- Implementing antibiotic policies;
- Education of staff, patients and visitors;
- Increasing space between beds and number of single rooms especially as new wards are built.

CHAPTER 7

he Kidney Patient Care Co-ordinators, at Beaumont Hospital, will provide you with practical and social support and advice, will liaise with you and your family regarding any particular difficulties which your illness presents, and will advise you accordingly. This encompasses a wide range of topics, from the different treatment options to practical help with housing, medical cards etc.

Once you have been diagnosed with End Stage Kidney Disease (ESKD), you will be referred to one of the Patient Care Co-ordinators either through the ward, the outpatients department, or your Consultant. Your allocated co-ordinator will then provide information on education, support, practical advice and any other assistance possible to help you in the transition to kidney replacement therapy (RRT).

EDUCATION

The Patient Care Co-ordinators, in conjunction with other members of the kidney team, run education sessions from time to time. These are specifically aimed at patients who are approaching ESKD and aim to answer the many questions that you will have at this time. They include information on the different treatments available for kidney disease such as haemodialysis, peritoneal dialysis or transplantation, and explain the options available to you, at this time. If you are unable to attend one of these sessions or, if there is no session planned at the time of your diagnosis, the Patient Care Coordinator will provide all the necessary information on a one-to-one basis.

PRACTICAL ADVICE

When you, or someone in your family, is diagnosed with a chronic illness such as kidney disease, the resulting non-medical financial burden can further complicate the stress involved, and it is important to become aware of the various assistance schemes available from the Department of Employment Affairs and Social Protection, the HSE and Local Authorities.

Not everyone is automatically entitled to financial assistance and some schemes are means tested. Specific individual circumstances are also taken into account; therefore, one person may qualify for assistance, while another, with the same illness, may not. You can only find out about your position by making an application and providing all the details required so that the relevant authority can determine your eligibility. Your Patient Care Co-ordinator can provide advice on your entitlements and, where applicable, provide a letter of support to accompany your application.

"Not everyone is automatically entitled to financial assistance, and some schemes are means tested."

The area of grants/allowances/benefits is a wide-ranging one that cannot be covered in detail in this publication. On the following pages is an overview of what financial assistance may be available and included at the end is a list of the relevant organisations to which you can apply for further information.

BENEFITS AND ENTITLEMENTS

There are three categories of eligibility for health services: medical card holders, GP visit card holders and non-medical card holders.

Medical card holders are entitled to free hospitalisation, GP services, most prescribed drugs and a range of other health servies. A medical card is means tested, based on the applicant's weekly income less PRSI. There is a prescription charge with your medical card.

Further information and a medical card application form can be obtained from your local health centre or can be downloaded from **www.citizensinformation.ie** or **www.hse.ie**

Hardship cases are dealt with on merit and special circumstances such as chronic illness can be taken into account. In cases of financial hardship, medical card holders may apply to the HSE office for assistance with the cost of on-going prescribed medical items not available under the

Medical Card Scheme. If you feel you need a medical card, do apply for one. A supporting letter may be obtained from your attending hospital on request.

GP visit card holders are entitled to free visits to their GP. This card is issued based on specific income guidelines. In some cases, where a person may have a chronic illness which involves regular GP visits, the HSE may grant the GP visit card even where their income is greater than the guidelines. Largely, the HSE will only consider these applications where an ongoing medical condition is causing, or likely to cause, undue financial hardship.

Non-medical card holders are liable for a Government Levy for in-patient stays. Non-medical card holders can avail of the Drugs Payment Scheme through their local pharmacy. Under this scheme, families (patients, their partners and dependant children) pay a fixed amount per month for prescribed medicine.

TAX RELIEF ON MEDICAL EXPENSES

You can claim tax relief on medical expenses you pay for yourself and on behalf of any other person. You can claim relief only if you cannot recover the expenses from any other source.

Further information is available on the Revenue Commissioner website **www.revenue.ie** Telephone: 1890 306706.

Tax relief is granted under specific headings in respect of certain expenses incurred by kidney patients. The amount of relief depends on where the patient receives care (for example, in hospital or at home). Home dialysis and CAPD patients may claim tax relief on items such as electricity, telephone, laundry and travel expenses.

DISABILITY ENTITLEMENTS

Patients who were employed, pre-dialysis, are fully encouraged to continue in their work/full-time education/training. However, there are cases where this is not an option and there is a range of entitlements to assist those who find themselves on a reduced income due to illness.

Illness Benefit is a short-term payment paid to insured people who are unfit for work due to illness. You will qualify if you:

- are under 66;
- are unfit for work due to illness;
- satisfy the PRSI contribution conditions.

Invalidity Pension is a long-term payment. To qualify, you must be permanently incapable of working. You must satisfy both PRSI and medical conditions. Since 1 December 2017, self-employed people who have paid sufficient class S PRSI contributions are entitled to Invalidity Pension.

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Disability Allowance is a long-term weekly allowance paid to people with a disability aged between 16 and 66. Your disability must be expected to last for at least one year and the allowance is subject to both medical suitability, means test and habitual residency test.

Both Invalidity Pension and Disability Allowance entitle you to a free travel pass and you may qualify for a companion free pass if you are unfit to travel alone. When you reach 66 years of age you no longer qualify for DA, but you are assessed for a State pension.

Working and Claiming a Disability Payment:

In certain circumstances, it may be possible to obtain a disability payment and work, provided the work is certified as being of a rehabilitative nature. People on Disability Allowance or Blind Persons' Pension may be allowed to retain their social welfare payment while working part-time (certain conditions apply). Written approval must be obtained from the Department of Employment Affairs and Social Protection. This may result in the withdrawal of a medical card.

Carer's Allowance is a payment to people on low incomes who are looking after a person who needs support because of age, disability or illness (including mental illness). This payment is meanstested. If a carer looks after more than one person, they may also be eligible for an additional payment of 50%. A carer may work outside the home for 15 hours per week provided this has first been approved with the Department of Employment Affairs and Social Protection. Any money earned, however, will be assessed as means in deciding the amount of allowance due.

There are other allowances available with the Carer's Allowance such as:-

- Free travel pass
- Household benefit package

Carer's Benefit: Those who leave the workforce to care for a person in need of full-time care and attention may be entitled to Carer's Benefit, which is based on PRSI contribution.

If you already receive a Social Welfare Payment you are not eligible for Carer's Allowance or Carer Benefit. You may claim either, in place of your Social Welfare Payment.

HEALTH SERVICE EXECUTIVE (HSE) PAYMENTS

Supplementary Welfare Allowance is an emergency payment for people without PRSI contributions. It is a basic minimum income to help bridge the gap while social welfare payments' applications are being processed. This payment is available from the local Community Welfare Officer.

Exceptional Needs Payment

An Exceptional Needs Payment is a single payment to help meet essential, once-off, exceptional expenditure, which a person could not reasonably be expected to meet out of their weekly income. For example, the payment can be

for bedding or cooking utensils for someone setting up a home for the first time, visiting relatives in hospital or prison, funeral costs or for clothing in exceptional circumstances.

Other HSE payments include:-

- Mobility Allowance
- Carers Support Grant
- Blind Welfare Allowance
- Motorised Transport Grant
- Back-to-School Clothing and Footwear Allowance.

These payments can be obtained by applying to your local HSE with supporting documentation.

SUPPLEMENTARY WELFARE ALLOWANCE SCHEME

The Supplementary Welfare Allowance Scheme is funded and legislated for by the Department of Employment Affairs and Social Protection. The payments made under this scheme include: **weekly Supplementary Welfare Allowance**, **Rent Supplement**, **Mortgage Interest Supplement**, **Diet and Heating Supplements**, **Back to School Clothing and Footwear Scheme** and exceptional needs payments for items such as buggies, clothing, etc. Your Patient Care Coordinator will advise on the relevant allowances for you and will write to the Department of Employment Affairs and Social Protection's representative to support your application as required.

For free information on your rights and entitlements contact: CITIZENS INFORMATION

Tel: 0761-07 4000 (Monday to Friday, 9am to 8pm)

or www.citizensinformation.ie

for your local Citizens Information Centre.

LOCAL AUTHORITY GRANTS

The Housing Adaptation Grant for people with a disability

This is a means tested grant to make a house suitable for a person with a disability. The grant can help you to make changes and adaptions to your home - for example, making it wheelchair accessible, extensions to create more space, adding ground floor bathroom or toilet or stair lift.

If you only require minor work you can apply for the Mobility Aids Grant Scheme. This ia a means tested grant to provide mobility equipment, e.g. the installation of grab-rails, a level access shower or a stair lift.

Housing Aid for Older People Scheme

This is a means-tested grant and is used to improve the condition of the older person's home. The type of work that is grant-aided includes structural repairs or improvements, rewiring repair or replacement of windows and doors, the provision of water, heating and sanitary services, cleaning and painting, or any other improvement work considered necessary.

In general, it is aimed at people 66 years of age and older, who are living in poor housing conditions. However, in cases of genuine hardship the local authority may assist people under age 66.

Apply to the housing department, of your local authority, for each of these schemes. You may need an Occupational Therapist (OT) to assess your daily living needs in support of your application.

SENIORS ALERT SCHEME

Grant support is available for the provision and installation of monitored personal alarms, which can be worn as a pendant or around your wrist like a watch. These alarms connect to a national helpline via a base unit installed in your home. When the alarm is activated, it automatically rings the helpline, which is open 24 hours a day all year round. A helpline operator talks to you in your home and decides whether to alert a local volunteer responder or, if necessary, the emergency services. Groups wishing to register for the scheme can register online at **pobal.ie**.

GOING HOME FROM HOSPITAL

The transition to kidney replacement therapy and/or transplantation is eased by the constant presence of medical and nursing staff while you remain in hospital. Once discharged, it is now time for you to adjust to this new way of life within the context of your own home and family life. There are many different ways that illness can affect your life. Maybe you are too unwell to do housework and need some home help. Perhaps you need meals-on-wheels. Whatever your requirements, your Patient Care Co-Ordinator will liaise with the relevant community services to ensure you have the appropriate support after your hospital stay.

PUBLIC HEALTH NURSE (PHN)

If you require any nursing care, following your discharge, you will be referred by the hospital to your Public Health Nurse, who is based in your local Health Centre. The Patient Care Co-Ordinator will ensure the PHN has all the necessary information required and will liaise with them with regard to your care.

HOME SUPPORT SERVICE

The Home Support service (previously know as Home Help Service or Home Care Package Scheme) provides help and support with every day tasks to older people living at home. These tasks may include help with:

- getting in and out of bed
- dressing and undressing
- personal care such as showering and shaving

The HSE wishes to support you to remain in your own home for as long as possible. As part of the

Home Support Service, HSE staff will consider what kind of support you might need. These supports will then be provided by HSE staff or by external service providers who have agreements with the HSE to provide this type of service on its behalf. If your home support is being delivered by an external provider, you will be asked to choose from a list of Approved Providers, which will be give to you by the HSE.

MEALS-ON-WHEELS

These are organised on a voluntary basis and a financial contribution



may be required. Your Patient Care Co-Ordinator or PHN will be able to ascertain if this service is available in your area and, if so, will make the necessary arrangements.

HSE LOCAL HEALTH OFFICES

Your local health office is your entry point to community health and personal social services. The wide range of services that are provided through local Health Offices and from Health Centres include general practitioner services, public health nursing, child health services, community welfare, chiropody, ophthalmic, speech therapy, social work, addiction counselling and treatment, physiotherapy, occupational therapy, psychiatric services and home help.

REPRODUCTIVE MATTERS

CHAPTER 8

SEXUAL ISSUES

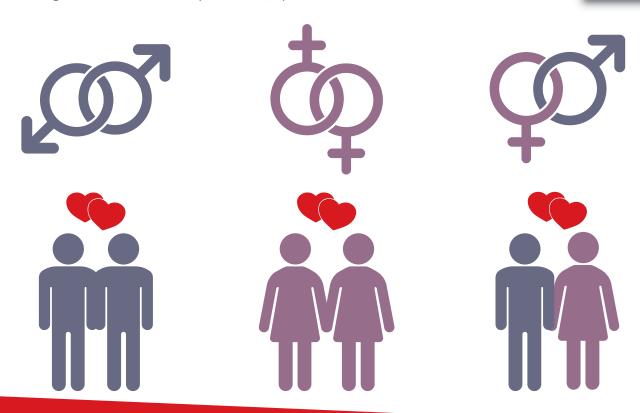
exual problems are common for men and women who suffer from kidney disease. Not only are emotional problems likely to occur as a result of the stress of the disease, but there are also a number of medical problems that can affect sexual function and fertility, both in men and women.

Emotional problems are common. Patients may find themselves going through a grieving process due to loss of kidney function, which may affect their independence, their job, and their role in the family. Some patients experience a change in body image. Dialysis can lead to lowered self-esteem, coupled with anger and depression, which can affect sexual function. The balance in the relationship may have changed – one seeing themselves as the carer and the other in a sick role. Couples need to communicate to one another their feelings and fears. Counselling can facilitate the exploration

of these feelings.

Contraception is important for people with kidney disease. Do not assume that because you have kidney disease, you cannot conceive a child. Most methods of contraception are suitable. methods such as condoms and diaphragms can be used. The coil can sometimes cause infection and heavy periods. The contraceptive pill has a tendency to raise blood pressure, sometimes one of the combined forms of oestrogen and progesterone prescribed. The morning after pill may be used in the usual manner.

Fertility levels vary during different stages of kidney disease. Women of childbearing age do not often get pregnant while on kidney replacement therapy as the treatment only replaces a small percentage of kidney function, which, in turn, can interfere with egg production.



For female patients whose menstrual cycle remains, however irregular, it is possible to conceive. Although pregnancy is uncommon, some women have given birth. Due to risks to the mother and the high rate of miscarriage, patients are normally advised to take precautions against pregnancy. Some women decide to delay pregnancy for one or two years after they have had a kidney transplant.

Men with kidney disease may have a reduced sperm count and may experience difficulties in fathering a child. After successful transplantation, sperm numbers generally rise. Men with kidney

disease can have a variety

of sexual problems.

"Men with kidney disease may have a reduced sperm count of sexual problems.

These include loss of libido and ejaculatory problems.

and may experience
difficulties in worrying and most common is impotence.
child." Impotence often has physical causes and is usually a combination of factors.

- Poor blood supply occurs as part of the natural ageing process and is common in older men. It is particularly common in men with diabetes and kidney disease.
- The testicles may produce less of the male hormone, testosterone.
- Some drug treatments can contribute to impotency, the biggest culprits being 'betablockers' such as atenolol, propranolol, metoprolol and bisoprolol.
- Tiredness can affect sexual performance. This can be caused by anaemia, under-dialysis or other medical problems e.g., heart problems.
- Psychological or relationship issues.

Treatment for impotence is commenced by checking out general health. Anaemia, if present, is corrected, treatment time may need to be extended

or drug treatment may be changed.

"For female
Viagra is commonly prescribed.

Patients whose
Trials show that about 8 out of 10

remains,
however irregular,
it is possible to
conceive."

menstrual cycle

Trials show that about 8 out of 10 men benefit with improvement in erectile performance. Patients with angina or heart problems should not take Viagra. It is advisable to consult your unit doctor to ensure that it is a safe option in your

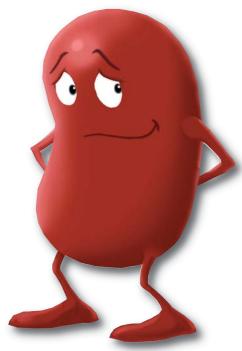
case.

There are many other interventions which are performed by specialist doctors e.g., urologists. A referral is sent from the unit doctor.

Counselling is recommended for emotional problems relating to impotence.

Many men, who have had difficulty fathering a child while undergoing kidney replacement therapy, have been successful in doing so following transplantation.

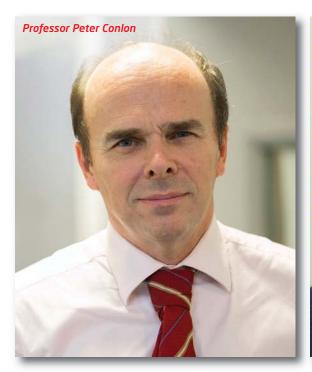
Communication is the most important factor in any relationship. Wherever there is a problem share your feelings and fears with your partner. Nursing and medical staff recognise that patients can have difficulties in sexual relationships and will gladly talk to you and your partner. Professional guidance can help - all that may be needed is a little reassurance.



CHAPTER 9

our kidney health care team has many years of experience treating people with kidney disease. If you have a question or are looking for information, ask any member of your healthcare team. If they can't help, they will refer you to someone who can. Do not hesitate to ask.

Because your kidney healthcare team is important to you, we describe, in this chapter, the roles and skills of each team member. This may help you to decide which people it may be beneficial to consult.





NEPHROLOGIST

A Nephrologist (or Consultant) is a doctor who specialises in kidney disease and who, together with you and the other members of the healthcare team, plans the best treatment for you.







The Nephrologist in a teaching hospital will typically have a team of non-consultant hospital doctors assisting them in your care. These doctors include the following:



RENAL REGISTRAR

A renal registrar will have completed at least 3 years of post-graduate medical education. They will be responsible for the supervision of the Senior House Officers and Interns. A registrar will typically be participating in a 4 to 5 year training programme on kidney diseases.

SENIOR HOUSE OFFICER

The senior house officer will be at least 1 year post-graduation from medical school and is training in the general aspects of hospital medicine.

INTERN

In the first year after graduation from medical school, all medical students must spend a year of training in the hospital to learn the fundamentals of hospital medicine

MEDICAL STUDENTS

Training hospitals will have many medical students. These students will talk to you about how the disease has affected you. They will also ask for permission to examine you. This is how they become a doctor themselves one day. On ward rounds frequently the Registrar or Consultant will ask the medical student to present their findings about you and your condition to the team. It is your choice to specify that you do not wish to see medical students if you are not feeling up to it.



NEPHROLOGY NURSE

Nephrology nurses work closely with you, your family and other team members. They will teach you and your family about your kidney disease and its treatment, and support you in the lifestyle changes that you may need to make. In some units, Clinical Nurse Specialists, Nurse Clinicians or Nurse Practitioners may also be part of the team. On each ward, a Clinical Nurse Manager will be responsible for the day-to-day management of the unit.



RENAL COUNSELLOR

Many kidney units in Ireland have access to a Counsellor who has specific experience in looking after patients with chronic kidney disease. Being diagnosed with kidney disease can be very difficult for you, your partner and your family. The Renal Counsellor will work closely with you and your family to help adjust to this new lifestyle.

STUDENT NURSES

Currently it takes four years at university to become a registered nurse. Student nurses spend much of this time rotating in different areas within the hospital. Student nurses will have different levels of experience, according to what year they are studying in university.

DIETICIAN

The dietician will instruct you on the proper food choices you may need to make as part of your treatment. Following consultation with your doctor and yourself, the dietician will then prepare a daily eating plan.



Olive McEnroe, Ruth O'Malley and Louise McSkeane Ambulatory Care Nurses

AMBULATORY CARE

You will meet the Ambulatory Care Nurses in the out-patient department. They will provide you with education and support and will make referrals to other members of the team, on your behalf. They will also co-ordinate the necessary tests if you are deemed suitable for a transplant.

PATIENT CARE CO-ORDINATOR (PCC)

Your Patient Care Co-Ordinator is available to provide supportive counselling to you and your

family. You may benefit from discussing emotional, financial, family or other concerns with your Patient Care Co-Ordinator as you try to understand and adjust to the changes that result from



Mary 1. Murphy,
Patient Care Co-ordinator

having kidney disease. They can also assist you with information about community resources and financial aid programs.



Ms Dilly Little, Transplant Surgeon

SURGEON

A surgeon is a doctor specifically trained in surgery. A surgeon will perform the operation to establish access for dialysis, or to transplant a kidney.



YOUR FAMILY DOCTOR (GP)

It is important that you continue to visit your family doctor. He, or she, knows you and your family best. Your kidney healthcare team will take excellent care of any problem associated with your kidney disease. However, your family doctor is best suited to provide preventative healthcare check-ups, such as pap smears and prostate examinations, and to look after any other healthcare needs. The kidney healthcare team will communicate with your GP on a regular basis.

TRANSPLANT CO-ORDINATORS

(L-R): Laura Lynch, Laura Austin, Andrea Fitzmaurice, Grainne Lambe, Emma Harte.



TRANSPLANT CO-ORDINATOR

The role of the co-ordinator is to serve as recipient co-ordinator for people awaiting kidney transplant. They facilitate and co-ordinate live kidney donation as well as co-ordinating donor referrals.



CARE ATTENDANTS

Work closely with nursing staff in providing your nursing care.

PHARMACIST

Works closely with members of the kidney healthcare team. They provide you with information, surrounding your medications, to ensure you have full understanding of your treatment.

HOUSEHOLD OPERATIVES

Responsible for household activities at ward level. They assist with stocking of equipment and supplies.

PORTERING STAFF

Assist by transporting patients to, and from procedures.

WARD CLERKS

Will ensure that the correct information is gathered in relation to:

- Medical Card Number
- Health Cover

DOMESTIC STAFF

Responsible for ward cleaning to ensure hygiene standards are maintained.

CATERING STAFF

Ensure that the food you receive is of a high standard. Also, they will deliver any special diets that your dietician has ordered for you.

STAYING HEALTHY

CHAPTER 10



aking part in pleasant leisure activities and maintaining a well-balanced lifestyle can go a long way toward helping you stay healthy, and have fun! It is important to take part in social activities, sports and recreation events, and other pasttimes that you, and other members of your family and friends, enjoy. You may need to make a few adjustments, but they will be well worth it.



EXERCISE AND SPORTS

Exercise is vitally important to both your physical and mental health. Staying physically fit will give you more energy. With more energy, you will feel like doing more things which will improve your outlook and speed your return to your usual lifestyle. Talk to your healthcare team about a suitable exercise or sports program before you begin (or resume) these activities.



There are many benefits to be enjoyed from exercise and these include:

- Improved physical functioning
- Better blood pressure
- Improved muscle strength
- Lower level of blood fats
- Better sleep
- Better control of body weight
- Reduced risk of heart disease
- Development of stronger bones
- Reduced stress and depression
- Meeting people
- Having fun.

If you are interested in getting and staying fit, then remember the **F.I.T.T** principle. Studies have shown that for it to be effective you must exercise:

- Frequently
- Reach a minimum Intensity
- Continue for a minimum length of Time
- Do an appropriate Type of exercise



To ensure that you get a positive effect, exercise at least three times a week. Unless you exercise this often, your physical conditioning will not improve. Spread the exercise over the week. Do not do it all on consecutive days. This gives your body time to recover and build up your muscles and energy for the next time.

Intensity

Unless you exercise hard enough, you are not going to get much benefit from it. There are different ways to measure intensity. One of the most common is by measuring your heart rate as you exercise. Talk to your doctor or someone who specialises in physical conditioning.

Time

The exercise must last at least 15 minutes to be effective. Any less, and your physical conditioning will not change. If you're not able to exercise continuously for that long, try doing it a little less vigorously, or pause for a couple of minutes to catch your breath, then keep going. The longer you keep it up, the more good it will do you.

Type

Some exercises are better than others. Some of the best are walking, swimming, and riding a bicycle. They get most of your muscles working and increase your heart rate to a healthy level. You can also adjust the intensity to suit your level of fitness. Best of all, they're fun to do.

Before you start a vigorous exercise program, check with your healthcare team-they can advise you on which exercises are best for you, and which ones, if any, you should avoid.



GENERAL HEALTH ADVICE



Eat a healthy diet, this means eating a wide variety of foods, in the correct amounts, to ensure good health. *Important points:*

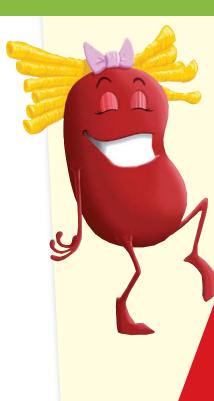
- Eat three main meals a day.
- Avoid fried and fatty foods.
- Eat more fruit and vegetables.
- Eat more fibre rich foods.
- Reduce sugary foods and sweetened drinks.

- Avoid adding salt to foods.
- Adhere to diet restrictions such as low salt, low cholesterol and diabetic diets according to doctors' or dieticians' instructions.
- Fluid restrictions may be necessary, please follow your doctor's advice regarding same.



Exercise has a positive affect on blood pressure, cholesterol levels and the functioning of the heart and lungs even if you do not lose weight.

Adopt a regular exercise pattern if one does not exist in your life. This is essential to maintain a healthy body weight and decrease the risk of obesity.



CLINIC APPOINTMENTS

Please attend all Out-Patient or other clinic appointments to ensure a continuous record of medical history. If you are unable to attend any scheduled appointments, please inform the department for further available dates.

"Please keep a
list of all current
medications with you
on all health visits as
it is important to have
an up-to-date
record."

MEDICATIONS

Please keep a list of your current medications with you on all health visits, as it is important to have an up-to-date record. Medications, not prescribed by your doctor, should not be taken.

Avoid herbal remedies without seeking medical advice, as some may interact with specific medications.

STRESS

Decreasing stress will improve your psychological health, which is essential to general health and well-being.

Further help may be sought from the kidney nurse counsellor or therapist.

SMOKING AND DRUGS

If you are currently smoking, try firstly to cut back and, over a period of time, to cease.

Smoking damages the lungs and puts you at risk of lung cancer.

Smoking tobacco, marijuana or other drug use is harmful to everyone.

Any form of 'recreational drugs', such as ecstasy, speed or cocaine, can have a serious

These drugs may also interact with your medications.

"Any form of 'recreational drugs' such as ecstasy, speed or cocaine can have a serious effect on your body and mind."



ALCOHOL INTAKE

Alcohol is high in sugars and calories. Excess alcohol can increase your triglyceride level and promote weight gain.

Use sugar free mixers such as diet minerals or slimline tonic.

Have 2-3 alcohol free days per week and, when taking alcohol, do not exceed the recommended limits:

(A) Men: 17 units per week.

(B) Women: 11 units per week.

The Irish Kidney
Association Renal
Support Centre is
located in the grounds
of Beaumont Hospital,
just 100 metres walk
from the main hospital
entrance, is open all year
round and provides free
accommodation for all its
residents, who include:



- Families of renal patients from outside Dublin.
 It is available to all renal families no matter what Dublin hospital their family member is attending.
- Renal patients who have to travel long distances to see their consultant as an outpatient may stay overnight when accommodation is available.
- The Centre arranges counselling service as required by outpatients and their families. The counselling service is located at Donor House.
- Preference for accommodation is given to families of patients receiving transplants and families of the seriously ill.



Thirteen
en-suite
bedrooms some of
which can sleep up to
four persons. All rooms are on
ground floor level and have satellite TV, hairdryer,
refrigerator and ironing facilities.

Comfortable sittingroom/dayroom with satellite TV. Fully fitted kitchen where meals can be prepared by residents. Complimentary tea and coffee is provided by the Association for residents and guests. Soft drinks and snacks are available from vending machine.

Laundry room with washing powder supplied. Parking for overnight residents only.

The Centre is owned and funded by the IKA. Donations from residents and fundraising initiatives are most welcome.

The Centre is open to residents all year round. Day facilities are available Monday to Friday from 8.30am to 4.30pm, Saturday and Sunday 12pm-4pm.

For further information contact:

IKA Renal Support Centre, Beaumont Hospital, Dublin D09 Y5R3.

Telephone: 353-1-837 3952. Out of hours (Emergency only): 087-416 9907

Email: renalcentre@ika. ie

GLOSSARY	
TERM	DEFINITION
AKI - Acute kidney injury ALSO CALLED AKD Acute kidney disease	A sudden loss of kidney function that is often reversible.
AVF - Arteriovenous fistula	Vascular access for dialysis; joining an artery and vein together.
Anaemia	A shortage of red blood cells in the blood. One of the functions of the kidneys includes EPO (erythropoietin) production. When the kidneys fail, EPO is not made leading to anaemia.
ANCA - (Anti-neutrophil cytoplasmic antibody)	A type of antibody that is associated with vasculitis conditions.
APD - Automated peritoneal dialysis	Also known as CCPD. This is a form of peritoneal dialysis which is carried out overnight.
Arteries	Blood vessels that carry blood from the heart to the rest of the body.
Blood Tests	A test that is used to measure many substances in the body to ensure they are within normal/safe range.
Blood Pressure (B/P)	The pressure that the blood exerts against the walls of the arteries as it flows through them.
CAPD - Continuous ambulatory peritoneal dialysis	Infusion of fluid into the peritoneum, prolonged dwell period and then drainage. See Book 2
Central Venous Catheter (CVC)	Also known as permcath. A catheter with two ports inserted into a major central vein for the purpose of haemodialysis.
Creatinine	A waste substance produced by the muscles when they are used. The higher the blood creatinine level, the greater the indication of kidney disease.
Chronic kidney disease (CRD)	Slow onset of kidney disease which is irreversible.
Dehydration	Insufficient water in the body to maintain normal function.
Dialysis (HD)	An artificial process which removes chemical substances and water from the blood by passing it through an artificial kidney.
End Stage Kidney Disease (ESKD)	When renal replacement therapy is required.
Erythropoietin (EPO)	Hormone involved in production of Red Blood Cells.
Fluid Overload	The body contains excess water. This occurs in kidney disease as one of the functions of the kidney is to remove excess fluid.

GLOSSARY continued	
TERM	DEFINITION
Haematuria	Blood in the urine.
Hepatitis	An infection of the liver. Can be passed on by blood contact.
Kidneys	Two bean-shaped body organs (the size of your fist) where urine is produced. Functions of the kidney include removal of toxic waste, removal of excess fluid, controls blood pressure, helps to produce red blood cells and helps to keep bones strong and healthy.
Nephron	Small filtering unit in the kidney, made up of blood vessels and tubules.
Oedema	A build up of fluid causing swelling, especially ankles and the lungs.
Oliguric	Passing low levels of urine.
Potassium	A mineral that is normally present in the blood. Too much or too little can cause complications.
Transplantation	The replacement of an organ that is not working in the body with another donor organ. For Kidney Transplantation see Book 3.

CONTRIBUTORS

We would like to extend special thanks to the following members of the Renal Team at Beaumont for their contribution to this book.

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Dr. Darren Pachaippan
Martin Ferguson

Sheila Donlon Olive McEnroe Ruth O'Malley Binu Vasu

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Also, to the patients and staff who took time to contribute to editing this book.

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St. Damien's Ward

Renal Day Care

Patient Care Co-Ordinators

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Prof. Conlon's Secretary

Dr. Magee's Secretary

Dr. Denton's Secretary

Prof. de Freitas's Secretary

Dr. O'Seaghdha's Secretary

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01-828 2751

01-809 2321

01-809 2747

01-797 4701

01-809 3080

01-809 3357

01-809 2567

01-852 8152

Useful Information Websites

BEAUMONT RENAL UNIT - www.beaumont.ie/kidneycentre

IRISH KIDNEY ASSOCIATION - www.ika.ie

IRISH HEALTH WEBSITE - www.irishhealth.com

AMERICAN ASSOCIATION OF KIDNEY PATIENTS - www.aakp.org

NATIONAL KIDNEY FOUNDATION USA - www.kidney.org

Kidney Dialysis Centres

Beaumont Hospital, Dublin

Tel: 01-809 3000

Beacon Renal, Sandyford, Dublin

Tel: 01-906 5628

Beacon Renal, Tallaght, Dublin

Tel: 01-906 5624

Beacon Renal, Drogheda, Dublin

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Cavan General Hospital

Tel: 049-437 6032

Cork University Hospital

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Daisyhill Hospital, Newry

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Tel: 056-777 2751

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Fresenius Northern Cross Dublin

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Mayo General Hospital

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Merlin Park Hospital, Galway

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Our Lady's Children's Hospital, Crumlin, Dublin

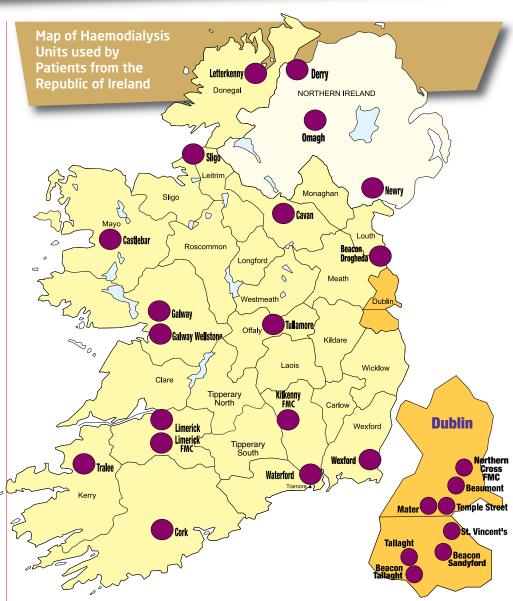
Tel: 01-409 6029/409 6948

Renal Care Centre, Wexford

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Sligo General Hospital

Tel: 071-917 4598



St. Vincents University Hospital, Dublin

Tel: 01-221 4427/221 3089

Tallaght Hospital, Dublin

Tel: 01-414 2358/414 2350

Temple Street Children's Hospital, Dublin

Tel: 01-878 4757

Tralee General Hospital

Tel: 066-718 4330

Tullamore Regional Hospital

Tel: 057-935 8743 /935 58740

Tyrone County Hospital, Omagh

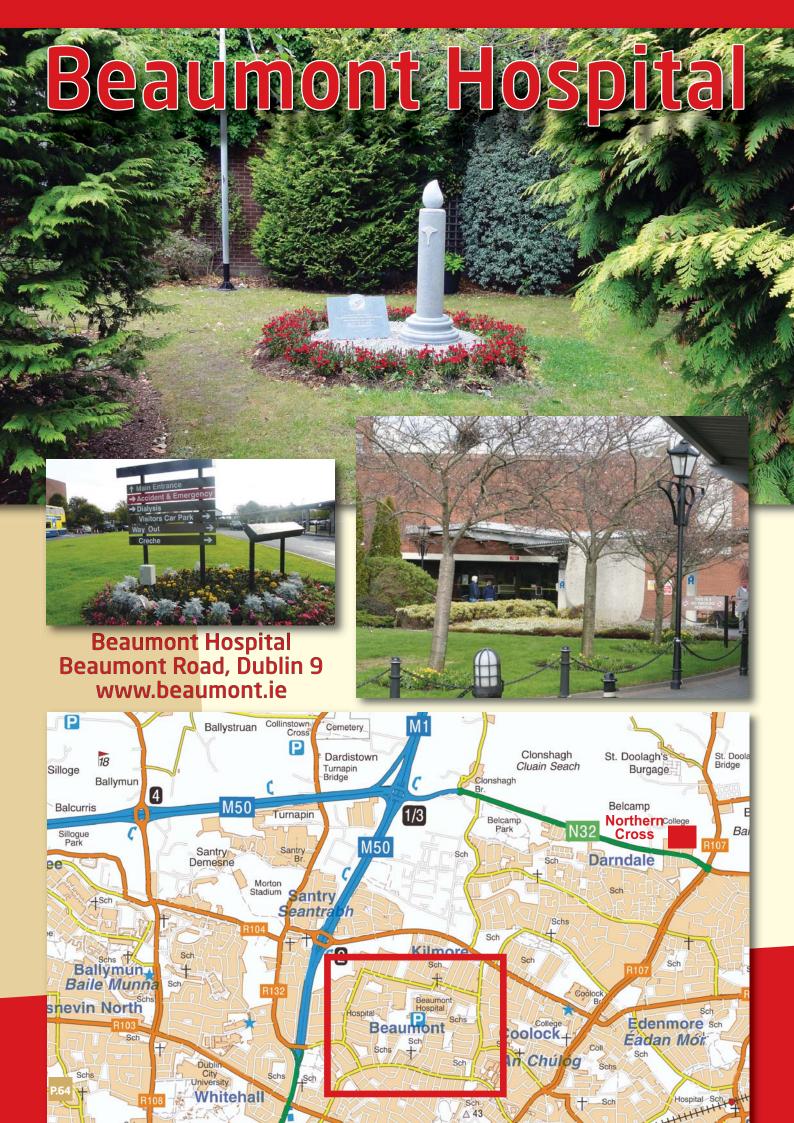
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Wellstone Ballybrit, Galway

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Beaumont Hospital Foundation

Established over twenty years ago, Beaumont Hospital Foundation is a registered charity that works to inspire charitable donations and promote support for its activities amongst patients, their families and friends, and locally and nationally from communities who access the services of Beaumont Hospital.

Funds raised by the Foundation are used to upgrade and purchase new, essential equipment that helps to ensure better care for patients and families.

As the National Renal Centre and home to the National Kidney Transplant Service, Beaumont Hospital is active in several areas of research to improve the lives of patients with kidney disease. Active areas of research include:

- The Irish Kidney Gene Project, which studies the genetic causes of kidney disease and operates a renal genetics clinic to help diagnose and treat those patients with hereditary kidney disease.
- A randomised trial of an app designed in Beaumont to improve fluid management in dialysis patients.
- Development of an app to improve potassium and phosphate control in dialysis patients.
- Development of an app to empower patients to better manage their kidney transplant.
- Observational studies of kidney transplant outcomes in Ireland.
- Observational studies of dialysis outcomes in Ireland.
- Clinical trials of novel biologic treatments for kidney disease.

All of these vital projects require funding in order to continue. If you would like to support a specific project, or make a general donation to Beaumont Renal research, you can do so online through our website or by phone on 01 - 809 2161.







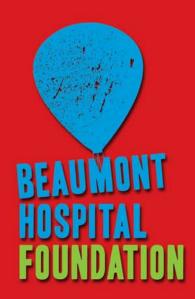




Beaumont Hospital Foundation, Beaumont Hospital, D 9

Tel: 01 - 809 2161 **Email:** hello@beaumont.ie

www.beaumont.ie/kidneycentre



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Beaumont Hospital, Dublin, D09 V2NO

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Web: www.beaumontfundraising.ie

CHARITY REGISTRATION NO: 11538



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