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**CURRENT FELLOWSHIP EXAM IN GENERAL SURGERY: A REVISION GUIDE**

**You Have to Pass Your FRCS Exam with Us**

A couple of surgeons performing surgery

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CAREFULLY SELECTED MCQ & EMQ QUESTIONS FROM PERVIOUS**12** YEARS EXAMS WITH ANSWER EXPLENATION AND UPDATED REVIEW

EDITED BY:

K.H. JAWAD -CONSULTANT SURGEON/FRCS-Ed

FIRST EDITION

January 1, 2021

KHALID SURGERY CLINIC

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The office location: Iraq/Baghdad 10053

Phone number: +9647706191881

Email: info@khalidsurgeryclinic.com

Website: https://www.khalidsurgeyclinic.com

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**Current fellowship exam in general surgery: A revision guide.**

The author: Dr. Khalid Hussun Jawad / consultant surgeon.

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**VOLUME ONE**

Dedication

Thanks, and gratitude to the woman who encouraged and helped me in the writing of this book.

To the woman who stayed up and was endured with me the hardness of this life all the time.

To the source of my inspiration and happiness in this world.

To my love and life, my dear wife MEMO I dedicates this book…

THE AUTHOR

**PREFACE**

Fellowship examination (FRCS) is greatly different from the membership examination (MRCS) which is taken care of basic and primary knowledge and sciences in surgery, and you will get a clinical diploma in surgery when you pass it, in contrast, the fellowship examination (FRCS) is more advancing. So, not only the basic and primary knowledge in surgery is required but usually, the situation is more complicated than the membership (MRCS) exam because you will become a consultant surgeon in the UK when you pass it. So, more knowledge is required, and more advanced decisions taken in surgical situations are needed.

FRCS exams are held by the four royal colleges of surgeons in the UK and divided into two sections. This book will help you to pass section one which is the written exam, and you will get a lot of information to fight in section two (clinical assessment).

Unlike the MRCS exam, the resources and books for the FRCS exam are limited, thus I decided to collect the frequent and recurrent questions for many years with other questions collected from the royal college of surgeon’s banks of questions and from different websites and designed to be a book of 15 chapters and regarded as a revision guide to everyone like to sit the exam. The questions and answers were arranged in such a way to create a great secret for learning ... So, this book contains a secret to pass your exam ... Every surgeon knows the practice, the practice, the practice is the secret of passing in the clinical assessment of any surgical exams. Repeating of information, and repeating, and repeating is the secret to pasting and keeping the surgical information in your brain and you will never forget them.

In this book, although the right answer is supported with explanation, they were updated to match more advanced surgical decision-making for current surgical practices.

I believe that this revision guide will make a great difference in your plan, and you will sit the examination with confidence and a comfortable minded

Thus, you have to pass your examination with this book.

Good luck

The Author

Dr. Khalid. H. Jawad.

**ACKNOWLEDGEMENTS**

I would like to extend my thanks and gratitude to the Royal College of Surgeons of Edinburgh for their providing continuous scientific support to me through their amazing website and allowing me free access to the private medical library and the college museum and providing me with all that is new in the science of surgery and sober scientific articles continuously.

I would like to thank the wonderful pastest website for providing me with a huge number of exam questions. I also extend my thanks and gratitude to the Blackwell’s Library in Edinburgh for the valuable books and medical references you provided me with.

Many thanks and gratitude to all my professors at the Royal College of Surgeons of Edinburgh who taught and trained me, thank you very much to all of them

All the love and appreciation to the professors in surgery, Professor Qalandar Hussein Kasnazan, and Professor Farouk Hassan Farag.

# INTRODUCTION

The British Fellowship in Surgery exam is one of the very important exams for surgeons to be accredited as consultants in surgical practice, and it is among the exams recognized in most countries of the world. It is held annually by the four royal colleges of surgeons in Britain and Ireland, and every doctor can participate in it after obtaining approval from the British Surgical Council, the participation and registration are done through the following link(https://www.jscfe.co.uk/). The FRCS exam is of two types, the first type is for doctors trained in and residing in Britain, and the other is called international for doctors from all over the world. And both types consist of two sections: writing exam and Clinical Assessment. The successful participant in the exam is granted the title of British Fellowship in surgery and the title of Clinical Consultant in Surgery. This book, although not the only one, is unique in its design and you can adopt it as a reference when you participate in the British Fellowship exam because it consists of repeating questions for many years, and if it indicates something, it only indicates the importance of these repeated questions for every surgeon who wants to be accredited as a consultant surgeon - the book consists of fifteen chapters in two volumes to facilitate carrying the book and reading. The chapters are arranged in a scientific way to simulate the recipient’s brain. The answers were supported by a sober scientific clarification and were renewed and updated according to the recent articles on each topic, which is what you are required to know as a consultant doctor and not what was mentioned, and you were restricted by the primary or basic books in surgery - because modernity is required to reach the level of the consultant surgeon.!

How do you read the book and pass the exam successfully?

You should read the book starting from the first chapter and sequentially until the last two chapters, which are the general and various topics in which the surgical knowledge you have reached for the purpose of participating in the exam will be evaluated. You should know a question differs from the other only in how it is formulated or with other information within the same content.!

Before you read, you should pay attention to the list of contents of each chapter in order to know how many topics were repeated, knowing that they were arranged in alphabetical order to make it easy to follow the repetition of each question and repetition each topic and in different places in the book to force the reader to follow up and to keep and paste the information in the mind - read the book once quickly and then read it once Others carefully - I will guarantee you success from the first trial.

When anyone participates in any exam in any subject and any topic in this life, the first thing that comes to the participant’s mind is what are the questions that can be asked? and what are the common and recurring questions? So, I have provided you in this book most of the questions asked by the committee designated questions - You should know that repeating the questions is not an indication of the inability of the competent committees to prepare the questions, but rather indicates the importance of that question and it will give you the required information to reach the level of the responsible consultant surgeon in your country or in Britain.

Good luck to you.

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**Miscellaneous**

1. **Theme:** Administration of local anaesthetics

|  |  |
| --- | --- |
| **A.** | 50ml of lignocaine 2% with 1 in 200,000 adrenaline |
| **B.** | 25ml of lignocaine 2% with 1 in 200,000 adrenaline |
| **C.** | 45ml 0.25% bupivicaine plain |
| **D.** | 45ml 0.5% bupivicaine with 1 in 200,000 adrenaline |
| **E.** | 25ml 1% lignocaine plain |
| **F.** | 15ml 1% lignocaine plain |
| **G.** | 30ml 1% lignocaine with 1 in 200,000 adrenaline |
| **H.** | 45ml 0.25% bupivicaine with 1 in 200,000 adrenaline |

Please select the most appropriate local anaesthetic choice for procedure and indication described. Each option may be used once, more than once or not at all.

|  |  |
| --- | --- |
| **1.** | An 80 kg 30 year old male undergoes a difficult appendicectomy via a lower midline abdominal incision. Prior to this event he was well.  http://d35pzpz673ryzz.cloudfront.net/css/small_tick.gif 45ml 0.25% bupivicaine plain  Wound infiltration is best performed using long acting local anaesthetic agent. |
| **2.** | A 50kg 65 year old women presents to the surgeons for excision of a lump from her thigh. She has longstanding depression for which she takes phenelzine. She is otherwise well.  **The correct answer is 15ml 1% lignocaine plain**  Adrenaline containing agents are contra indicated when MAOI drugs are present as they may precipitate a hypertensive crisis. |
| **3.** | A 50 kg 60 year old women presents to the surgeons for excision of lump from her back under local anaesthesia. She has longstanding depression for which she takes venlafaxine. She is otherwise well.  **The correct answer is 30ml 1% lignocaine with 1 in 200,000 adrenaline**  Surgery on the back is often bloody and use of an adrenaline containing solution can be beneficial. SNRI's are not a contra indication to the use of adrenaline containing agents. |

**Local anaesthetic agents**

**Lidocaine**

* An amide
* Local anaesthetic and a less commonly used antiarrhythmic (affects Na channels in the axon)
* Hepatic metabolism, protein bound, renally excreted
* Toxicity: due to IV or excess administration. Increased risk if liver dysfunction or low protein states. Note acidosis causes lidocaine to detach from protein binding.
* Drug interactions: Beta blockers, ciprofloxacin, phenytoin
* Features of toxicity: Initial CNS over activity then depression as lidocaine initially blocks inhibitory pathways then blocks both inhibitory and activating pathways. Cardiac arrhythmias.
* Increased doses may be used when combined with adrenaline to limit systemic absorption.

**Cocaine**

* Pure cocaine is a salt, usually cocaine hydrochloride. It is supplied for local anaesthetic purposes as a paste.
* It is supplied for clinical use in concentrations of 4 and 10%. It may be applied topically to the nasal mucosa. It has a rapid onset of action and has the additional advantage of causing marked vasoconstriction.
* It is lipophillic and will readily cross the blood brain barrier. Its systemic effects also include cardiac arrhythmias and tachycardia.
* Apart from its limited use in ENT surgery it is otherwise used rarely in mainstream surgical practice.

**Bupivacaine**

* Bupivacaine binds to the intracellular portion of sodium channels and blocks sodium influx into nerve cells, which prevents depolarization.
* It has a much longer duration of action than lignocaine and this is of use in that it may be used for topical wound infiltration at the conclusion of surgical procedures with long duration analgesic effect.
* It is cardiotoxic and is therefore contra indicated in regional blockage in case the tourniquet fails.
* Levobupivicaine (Chirocaine) is less cardiotoxic and causes less vasodilation.

**Prilocaine**

* Similar mechanism of action to other local anaesthetic agents. However, it is far less cardiotoxic and is therefore the agent of choice for intravenous regional anaesthesia e.g. Biers Block.

All local anaesthetic agents dissociate in tissues and this contributes to their therapeutic effect. The dissociation constant shifts in tissues that are acidic e.g. where an abscess is present, and this reduces the efficacy.  
  
**Doses of local anaesthetics**

| **Agent** | **Dose plain** | **Dose with adrenaline** |
| --- | --- | --- |
| Lignocaine | 3mg/Kg | 7mg/Kg |
| Bupivacaine | 2mg/Kg | 2mg/Kg |
| Prilocaine | 6mg/Kg | 9mg/Kg |

These are a guide only as actual doses depend on site of administration, tissue vascularity and co-morbidities.

**Maximum total local anaesthetic doses**

* Lignocaine 1% plain - 3mg/ Kg - 200mg (20ml)
* Lignocaine 1% with 1 in 200,000 adrenaline - 7mg/Kg - 500mg (50ml)
* Bupivicaine 0.5% - 2mg/kg- 150mg (30ml)

*Maximum doses are based on ideal body weight*

**Effects of adrenaline**  
Adrenaline may be added to local anaesthetic drugs. It prolongs the duration of action at the site of injection and permits usage of higher doses (see above). It is contra indicated in patients taking MAOI's or tricyclic antidepressants. The toxicity of bupivacaine is related to protein binding and addition of adrenaline to this drug does not permit increases in the total dose of bupivacaine, in contrast to the situation with lignocaine.

1. **Botulinum toxin is a popular therapeutic agent. By what mechanism does it exert its effects?**

|  |  |
| --- | --- |
| A | Inhibits release of acetylcholine into the neuronal synapse |
| B | Blockade of the post synaptic acetylcholine receptor |
| C | Increased hydrolysis of acetylcholine within the synapse |
| D | Release of sterically altered acetylcholine into the synapse |
| E | Increased re-uptake from within the synapse |

**Botulinum toxin**

Botulinum toxins are produced by the bacterium *Clostridium botulinum*. The toxin is a two chain polypeptide. One of the chains is a protease that is able to attack fusion proteins at the neuromuscular junction preventing vesicles from anchoring to the membrane to release acetylcholine. By inhibiting acetylcholine release, the toxin interferes with nerve impulses and causes flaccid (sagging) paralysis of muscles in botulism, as opposed to the spastic paralysis seen in tetanus.   
Botulinum toxin A is the agent used therapeutically and small doses are typically administered to the desired area. The doses are small and so in some cases the effects of treatment are transient and will wear off with time. Repeated dosing can result in more enduring muscular paralysis.

1. **What is the effect of increasing the confidence limit from 95% to 99%?**

|  |  |
| --- | --- |
| A | Confidence interval widens |
| B | Confidence interval remains unchanged |
| C | Confidence interval narrows |
| D | The study data becomes less accurate |
| E | None of the above |

The confidence limit is expressed as a percentage, usually 95%. Increasing it will increase the range of values. For example if the 95% confidence interval was between 50 and 55 and the confidence limit were increased to 99% then the value range would widen (say from 48 to 57). Conversely decreasing the confidence limit would decrease the range of values (from 52 to 54, for example).

Confidence intervals

A 95% confidence interval is often interpreted as indicating a range within which we can be 95% certain that the true effect lies. This statement is a loose interpretation, but is useful as a rough guide. The strictly-correct interpretation of a confidence interval is based on the hypothetical notion of considering the results that would be obtained if the study were repeated many times. If a study were repeated infinitely often, and on each occasion a 95% confidence interval calculated, then ninety five percent of these intervals would contain the true effect.

1. **Theme:** Antimicrobial prophylaxis for gastrointestinal endoscopy

|  |  |
| --- | --- |
| **A.** | Intravenous teicoplanin |
| **B.** | Oral ciprofloxacin |
| **C.** | No antimicrobial prophylaxis |
| **D.** | Intravenous co-amoxylav |
| **E.** | Oral penicillin V |
| **F.** | Intravenous tazocin |

Please select the most appropriate antimicrobial prophylaxis for the following individuals who are undergoing gastrointestinal endoscopy. Each option should be used once, more than once or not at all.

|  |  |
| --- | --- |
| **1.** | A 38 year old man with primary sclerosing cholangitis needs an ERCP and biliary decompression.   **The correct answer is Oral ciprofloxacin**  PSC decompression is technically challenging and seldom achieved at the first attempt. Antimicrobial cover is recommended in such cases and ciprofloxacin is the agent of choice. |
| **2.** | A 45 year old women has mitral regurgitation and is due to undergo a colonoscopy.  http://d35pzpz673ryzz.cloudfront.net/css/small_tick.gif No antimicrobial prophylaxis  Antibiotics are not routinely indicated for prophylaxis for routine cases. |
| **3.** | A 38 year old lady is due to undergo an upper GI endoscopy with duodenal biopsies. She has a mechanical aortic valve.  **The correct answer is No antimicrobial prophylaxis**  Again, antibiotics are no longer routinely administered. |

**Antimicrobial cover for gastrointestinal endoscopy**

* Routine antibiotics are not indicated for diagnostic upper and lower GI endoscopy
* Antimicrobial prophylaxis is not longer advocated for those individuals with underlying cardiac disease undergoing upper and lower GI endoscopy
* Prophylaxis is indicated in some therapeutic procedures (in all patients) including: ERCP for cholangitis, PEG insertion, PEC insertion, endoscopic FNA

1. **Which of the following is a major component of cryoprecipitate?**

|  |  |
| --- | --- |
| A | Protein C |
| B | Protein S |
| C | Factor V |
| D | Factor VIII |
| E | Factor IX |

**Cryoprecipitate**

* Blood product made from plasma
* Usually transfused as 6 unit pool
* Indications include massive haemorrhage and uncontrolled bleeding due to haemophilia

**Composition**

| **Agent** | **Quantity** |
| --- | --- |
| Factor VIII | 100IU |
| Fibrinogen | 250mg |
| von Willebrand factor | Variable |
| Factor XIII | Variable |

1. **An 80 year old lady is investigated in the pre operative clinic and found to have severe aortic stenosis. What, if any, is the main peri operative concern?**

|  |  |
| --- | --- |
| A | They cannot adjust their heart rate |
| B | They may have ventricular hypertrophy |
| C | The patient cannot increase their cardiac output |
| D | They are more prone to arrhythmias |
| E | There is no concern |

Patients with aortic stenosis are a major perioperative concern. They may have ventricular hypertrophy and this can result in relative myocardial ischaemia and increase the risk of arrhymias. However, the main concern is that they cannot increase their cardiac output particularly if vasodilation occurs.

**Aortic stenosis**

* Narrowing of the aortic valve
* May occur as a result of rheumatic fever or with aging and calcific changes
* Congenitally may occur earlier due to calcification of a bicuspid aortic valve (1-2% of population)
* Symptoms include exertional angina and syncope
* Where the condition is suspected, trans thoracic echocardiography is the investigation of choice

**Severity**

| **Degree** | **Mean gradient (mmHg)** | **Aortic valve area (cm2)** |
| --- | --- | --- |
| Mild | <25 | >1.5 |
| Moderate | 25-40 | 1.0-1.5 |
| Severe | >40 | <1 |

**Treatment**  
Either transcatheter or open aortic valve replacement

1. **A 28 year old lady presents with a pigmented lesion on her calf. Excisional biopsy confirms a diagnosis of melanoma measuring 1cm in diameter with a Breslow thickness of 0.5mm. The lesion is close <1 mm to all resection margins. Which of the following surgical resection margins is acceptable for this lesion?**

|  |  |
| --- | --- |
| A | 2 cm |
| B | 0.5 cm |
| C | 3 cm |
| D | 5 cm |
| E | 1 cm |

**Malignant melanoma**

| **The main diagnostic features (major criteria):**   * **Change in size** * **Change in shape** * **Change in colour** | **Secondary features (minor criteria)**   * **Diameter >6mm** * **Inflammation** * **Oozing or bleeding** * **Altered sensation** |
| --- | --- |

**Treatment**

* Suspicious lesions should undergo excision biopsy. The lesion should be removed completely as incision biopsy can make subsequent histopathological assessment difficult.
* Once the diagnosis is confirmed the pathology report should be reviewed to determine whether further re-excision of margins is required (see below):

**Margins of excision-Related to Breslow thickness**

|  |  |
| --- | --- |
| **Lesions 0-1mm thick** | 1cm |
| **Lesions 1-2mm thick** | 1- 2cm (Depending upon site and pathological features) |
| **Lesions 2-4mm thick** | 2-3 cm (Depending upon site and pathological features) |
| **Lesions >4 mm thick** | 3cm |

Marsden J *et al*. Revised UK guidelines for management of Melanoma. *Br J Dermatol* 2010 **163**:238-256.  
  
Further treatments such as sentinel lymph node mapping, isolated limb perfusion and block dissection of regional lymph node groups should be selectively applied.

1. **Recall bias is most commonly associated with which study design?**

|  |  |
| --- | --- |
| A | Cohort study |
| B | Cross sectional study |
| C | Case control study |
| D | Randomised control trial |
| E | Blinded randomised control trial |

**Recall Bias**

Recall bias represents a major threat to the internal validity of studies using self-reported data. It arises with the tendency of subjects to report past events in a manner that is different between the two study groups. This pattern of recall errors can lead to differential misclassification of the related variable among study subjects with a subsequent distortion of measure of association in any direction from the null, depending on the magnitude and direction of the bias. Although recall bias has largely been viewed as a common concern in case-control studies, it also has been documented as an issue in some prospective cohort and randomized controlled trial designs.

1. **A 12 year old child is admitted with a 12 hour history of colicky right upper quadrant pain. On examination the child is afebrile and is jaundiced. The abdomen is soft and non tender at the time of examination. What is the most likely cause?**

|  |  |
| --- | --- |
| A | Infectious hepatitis |
| B | Acute cholecystitis |
| C | Hereditary spherocytosis |
| D | Gilberts syndrome |
| E | Crigler najjar syndrome |

The child is most likely to have hereditary spherocytosis. In these individuals there may be disease flares precipitated by acute illness. They form small pigment stones. These may cause biliary colic and some may require cholecystectomy.

Hereditary spherocytosis

Most common disorder of the red cell membrane, it has an incidence of 1 in 5000. The abnormally shaped erythrocytes are prone to splenic sequestration and destruction. This can result in hyperbilirubinaemia, jaundice and splenomegaly. In older patients an intercurrent illness may increase the rate of red cell destruction resulting in more acute symptoms.   
Severe cases may benefit from splenectomy.

1. **Theme:** Choice of statistical methods

|  |  |
| --- | --- |
| **A.** | Students T Test |
| **B.** | Fishers exact test |
| **C.** | Mann Whitney U test |
| **D.** | Chi Squared test |
| **E.** | Spearmans rank test |

Please select the most appropriate statistical method for the situation described. Each option may be used once , more than once or not at all.

|  |  |
| --- | --- |
| **1.** | Concerns are raised about the high frequency of admissions to intensive care units following incisional hernia repairs. It is necessary to determine whether the rates of admission are significant.  **The correct answer is Chi Squared test**  This is dichotomous data (i.e. admitted or not admitted). Comparison between two groups will be needed (probably from different centres). Numbers are likely to be large and therefore the Chi Squared the most effective test. |
| **2.** | Researchers wish to determine whether there is an association between body weight and anastomotic dehisence following low anterior resection.   **The correct answer is Students T Test**  Body weight is likely to follow the normal distribution (provided the sample size is large enough). Therefore a T test is a suitable method for analysis. |
| **3.** | Surgeons wish to determine whether there is a relationship between operating time and lymph node yield in oesophageal surgery.   **The correct answer is Spearmans rank test**  This is testing a relationship between two numerical variables. Data is non parametric and a Spearmans rank test the most appropriate method. |

**Statistics**

**Data types**  
Accurately classifying the data you seek to obtain is the first step in undertaking formal data analysis.

| **Title** | **Description** |
| --- | --- |
| Nominal | Numbers are assigned to data that has no underling numerical value (e.g. marital status) |
| Ordinal | Has numbers that can be assigned to a natural underlying order (e.g. tumour grades) |
| Discrete | Data has a discrete numerical value, that has to be a whole number (e.g. number of deaths) |
| Continuous | Data has a numerical value that may not be a whole number and often reflects a direct measurement (e.g. weight) |

Knowing the data types allows us to direct the appropriate analysis. This is often conveniently achieved by plotting it on a graph. Where the data has a categorical nature, a histogram is often a useful starting point. Other types of data, particularly direct measurements, may be plotted as single data points. If we take the weight example from above then plotting a large number of data points may allow us to numerically determine the spread of the data. In particular whether it fits the normal distribution. Remember that if the mean, median and mode overlap numerically then the data will be normally distributed.   
  
**Parametric vs Non parametric**  
Parametric methods of data analysis assume that the underlying data set has a normal distribution. Non parametric methods do not make assumptions about the nature of the underlying data. 

| **Parametric tests** | **Non parametric tests** |
| --- | --- |
| T Test | Mann Whitney U |
| Paired T Test | Chi Squared |
|  | Spearmans Rank Correlation |
|  | Wilcoxon signed rank test |

*There are many others*  
  
**Types of test**

| **Test type** | **Features** |
| --- | --- |
| T Test | Direct comparison of data sets which are normally distributed |
| Mann Whitney U | Ranked method for non parametric data |
| Wilcoxon matched pairs/ signed rank | Analog of the paired T Test, data must be interval, data based on magnitude of differences |
| Spearmans Rank Correlation | Statistical dependence between 2 variables. May be used for continuous or discrete variables |
| Chi Squared test | Test of association between two qualitative variables,valid if 80% expected frequencies exceed 5 or all exceed 1. Fishers exact test may be used for small samples |

1. **Which of the following statements relating to the Chi Squared test is true?**

|  |  |
| --- | --- |
| A | The test makes comparisons between groups |
| B | The test is more useful if small numbers are present |
| C | The test requires prior calculation of the potency ratio |
| D | A probability of p <0.1 indicates statistical significance |
| E | The test compares actual values rather than proportions or number of occurrences |

Note that where numbers are less than 40, Yates correction may be required, this can reduce the value of Chi Square. The numbers used in Chi Squared tests are proportions rather than actual values.

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| Chi Squared test | Test of association between two qualitative variables,valid if 80% expected frequencies exceed 5 or all exceed 1. Fishers exact test may be used for small samples |

1. **Which of the following has the greatest impact on the positive predictive value of a test?**

|  |  |
| --- | --- |
| A | Prevalence |
| B | Subjects who are true negatives |
| C | Specificity |
| D | Relative risk |
| E | None of the above |

The positive predictive value (PPV) is the probability that an individual with a positive screening result has the disease. The sensitivity is the probability that an individual with the disease is screened positive and the specificity is the probability that an individual without the disease is screened negative.  
Its value depends upon the prevalence of the condition being tested for and the sensitivity of the test used.  
It may be calculated by dividing the number of true positives by the number of true positives and the number of false positives.

**Positive predictive values**

**Screening tests**

* Sensitivity: proportion of true positives identified by a test
* Specificity: proportion of true negatives correctly identified by a test
* Positive predictive value: proportion of those who have a positive test who actually have the disease
* Negative predictive value: proportion of those who test negative who do not have the disease

Predictive values are dependent on the prevalence

* Likelihood ratio for a positive test result = sensitivity/(1-specificity)
* Likelihood ratio for a negative test result = (1-sensitivity)/specificity

Likelihood ratios are not prevalence dependent

1. **Theme:** ASA grading

|  |  |
| --- | --- |
| **A.** | ASA I |
| **B.** | ASA II |
| **C.** | ASA III |
| **D.** | ASA IV |
| **E.** | ASA V |

The American society of anaesthesiologists physical status scoring system is a popular method for stratifying patients physical status. Please select the most appropriate ASA grade for each of the following scenarios. Each option may be used once, more than once or not at all.

|  |  |
| --- | --- |
| **1.** | A 66 year old man is admitted following a collapse whilst waiting for a bus. Clinical examination confirms a ruptured abdominal aortic aneurysm. He is moribund and hypotensive  http://d35pzpz673ryzz.cloudfront.net/css/small_tick.gif ASA V  Patients who are moribund and will not survive without surgery are graded as ASA 5. |
| **2.** | A 23 year old man with a 4cm lipoma on his flank is due to have this removed as a daycase. He is otherwise well.  http://d35pzpz673ryzz.cloudfront.net/css/small_tick.gif ASA I  Absence of co-morbidities and small procedure with no systemic compromise will equate to an ASA score of 1. |
| **3.** | A 72 year old man is due to undergo an inguinal hernia repair. He suffers from COPD and has an exercise tolerance of 10 yards. He also has pitting oedema to the thighs.  **The correct answer is ASA IV**  Severe systemic disease of this nature is a constant threat to life. Especially as he also has evidence of cardiac failure. |

American Society of anesthesiologists physical status scoring system (ASA**)**

| **ASA grade** | **Description** |
| --- | --- |
| 1 | No organic physiological, biochemical or psychiatric disturbance. The surgical pathology is localised and has not invoked systemic disturbance. |
| 2 | Mild or moderate systemic disruption caused either by the surgical disease process or though underlying pre-existing disease |
| 3 | Severe systemic disruption caused either by the surgical pathology or pre-existing disease |
| 4 | Patient has severe systemic disease that is a constant threat to life |
| 5 | A patient who is moribund and will not survive without surgery |

1. **In 2010 the journal Nature was stated to have an impact factor of 30.98. Which of the following is not needed to derive this information?**

|  |  |
| --- | --- |
| A | The total number of citable articles published by Nature in 2009 |
| B | The total number of citable articles published by Nature in 2008 |
| C | The number of times that articles published by Nature in 2008 were cited by articles in indexed journals in 2010 |
| D | The number of citable articles published in 2007 |
| E | The number of times that articles published by Nature in 2009 were cited by articles in indexed journals in 2010 |

The number of citable articles published by the journal in the preceding 2 years prior to the year in which the impact factor is quoted is needed.

**Impact factor**

The impact factor of an academic journal is a measure reflecting the average number of citations to recent articles published in the journal. It is frequently used as a proxy for the relative importance of a journal within its field, with journals with higher impact factors deemed to be more important than those with lower ones. The impact factor was devised by Eugene Garfield, the founder of the Institute for Scientific Information. Impact factors are calculated yearly starting from 1975 for those journals that are indexed in the Journal Citation Reports.  
  
**Calculation**  
In a given year, the impact factor of a journal is the average number of citations received per paper published in that journal during the two preceding years. For example, if a journal has an impact factor of 10 in 2007, then its papers published in 2005 and 2006 received 10 citations each on average in 2007. The 2007 impact factor of a journal would be calculated as follows:  
  
A = the number of times that articles published in that journal in 2005 and 2006, were cited by articles in indexed journals during 2007.  
B = the total number of "citable items" published by that journal in 2005 and 2006. ("Citable items" are usually articles, reviews, proceedings, or notes; not editorials or letters to the editor.)  
  
2007 impact factor = A/B.  
  
(Note that 2007 impact factors are actually published in 2008; they cannot be calculated until all of the 2007 publications have been processed by the indexing agency.)  
  
New journals, which are indexed from their first published issue, will receive an impact factor after two years of indexing; in this case, the citations to the year prior to Volume 1, and the number of articles published in the year prior to Volume 1 are known zero values. Journals that are indexed starting with a volume other than the first volume will not get an impact factor until they have been indexed for three years. Annuals and other irregular publications sometimes publish no items in a particular year, affecting the count. The impact factor relates to a specific time period; it is possible to calculate it for any desired period, and the Journal Citation Reports (JCR) also includes a five-year impact factor.

1. **A 56 year old man presents with symptoms of neuropathic facial pain and some weakness of the muscles of facial expression on the right side. On examination he has a hard mass approximately 6cm anterior to the right external auditory meatus. What is the most likely diagnosis?**

|  |  |
| --- | --- |
| A | Adenoid cystic carcinoma |
| B | Lymphoma |
| C | Adenocarcinoma |
| D | Pleomorphic adenoma |
| E | Mucoepidermoid carcinoma |

The patient is most likely to have a malignant lesion within the parotid. Of the malignancies listed; adenoid cystic carcinoma has the greatest tendency to perineural invasion.

Please rate this question:

**Parotid gland disease**

* Most parotid neoplasms (80%) are benign lesions
* Most commonly present with painless mass in cheek region
* Up to 30% may present with pain, when this is associated with a discrete mass lesion in the parotid it usually indicates perineural invasion.
* Perineural invasion is very unlikely to occur in association with benign lesions
* 80% of patients with facial nerve weakness caused by parotid malignancies will have nodal metastasis and a 5 year survival of 25%

**Types of malignancy**

| **Type of lesion** | **Features** |
| --- | --- |
| **Mucoepidermoid carcinoma** | 30% of all parotid malignancies Usually low potential for local invasiveness and metastasis (depends mainly on grade) |
| **Adenoid cystic carcinoma** | Unpredictable growth pattern Tendency for perineural spread Nerve growth may display skip lesions resulting in incomplete excision Distant metastasis more common (visceral rather than nodal spread) 5 year survival 35% |
| **Mixed tumours** | Often a malignancy occurring in a previously benign parotid lesion |
| **Acinic cell carcinoma** | Intermediate grade malignancy May show perineural invasion Low potential for distant metastasis 5 year survival 80% |
| **Adenocarcinoma** | Develops from secretory portion of gland Risk of regional nodal and distant metastasis 5 year survival depends upon stage at presentation, may be up to 75% with small lesions with no nodal involvement |
| **Lymphoma** | Large rubbery lesion, may occur in association with Warthins tumours Diagnosis should be based on regional nodal biopsy rather than parotid resection Treatment is with chemotherapy (and radiotherapy) |

1. **Which of the processes listed below is least likely to reduce the possibility of systematic errors arising from a clinical study?**

|  |  |
| --- | --- |
| A | Logarithmic transformation |
| B | Stratification |
| C | Randomisation |
| D | Restricted entry criteria |
| E | Logistic regression |

Logarithmic transformation is used to compare data sets that have different variances or are non normally distributed.

**Statistics**

**Data types**  
Accurately classifying the data you seek to obtain is the first step in undertaking formal data analysis.

| **Title** | **Description** |
| --- | --- |
| Nominal | Numbers are assigned to data that has no underling numerical value (e.g. marital status) |
| Ordinal | Has numbers that can be assigned to a natural underlying order (e.g. tumour grades) |
| Discrete | Data has a discrete numerical value, that has to be a whole number (e.g. number of deaths) |
| Continuous | Data has a numerical value that may not be a whole number and often reflects a direct measurement (e.g. weight) |

Knowing the data types allows us to direct the appropriate analysis. This is often conveniently achieved by plotting it on a graph. Where the data has a categorical nature, a histogram is often a useful starting point. Other types of data, particularly direct measurements, may be plotted as single data points. If we take the weight example from above then plotting a large number of data points may allow us to numerically determine the spread of the data. In particular whether it fits the normal distribution. Remember that if the mean, median and mode overlap numerically then the data will be normally distributed.   
  
**Parametric vs Non parametric**  
Parametric methods of data analysis assume that the underlying data set has a normal distribution. Non parametric methods do not make assumptions about the nature of the underlying data. 

| **Parametric tests** | **Non parametric tests** |
| --- | --- |
| T Test | Mann Whitney U |
| Paired T Test | Chi Squared |
|  | Spearmans Rank Correlation |
|  | Wilcoxon signed rank test |

*There are many others*  
  
**Types of test**

| **Test type** | **Features** |
| --- | --- |
| T Test | Direct comparison of data sets which are normally distributed |
| Mann Whitney U | Ranked method for non parametric data |
| Wilcoxon matched pairs/ signed rank | Analog of the paired T Test, data must be interval, data based on magnitude of differences |
| Spearmans Rank Correlation | Statistical dependence between 2 variables. May be used for continuous or discrete variables |
| Chi Squared test | Test of association between two qualitative variables,valid if 80% expected frequencies exceed 5 or all exceed 1. Fishers exact test may be used for small samples |

1. **Which of the following is a not a diagnostic criteria for brain death?**

|  |  |
| --- | --- |
| A | No response to sound |
| B | No corneal reflex |
| C | Absent vestibular-cochlear reflex |
| D | No response to supra orbital pressure |
| E | No cough reflex with bronchial stimulation |

**Brain death**

**Conditions for brainstem death testing**  
There must be an identifiable pathology causing irremediable brain damage. This may be intra - or extra - cranial.  
  
2. The patient must be deeply unconscious.  
a. Hypothermia must be excluded as the cause of unconsciousness and the patients core temperature should be over 34oC.  
b. There should be no evidence that the patients state is due to depressant drugs. This refers to narcotics, hypnotics and tranquillisers as well as neuromuscular blocking drugs. A  
careful drug history is required, whilst drug levels and antagonists may need to be used.  
c. Potentially reversible circulatory, metabolic and endocrine disturbances must have been  
excluded as the cause of the continuing unconsciousness. Some of these disturbances may occur as a result of the condition rather than the cause and these do not preclude the diagnosis of brain stem death.  
  
3. The patient must be apnoeic, needing mechanical ventilation. This condition must not be secondary to the effect of sedative drugs of neuromuscular blockade. This may require testing with a nerve stimulator to show intact neuromuscular transmission. Alternatively, demonstration of tendon reflexes can also demonstrate intact transmission  
  
**Criteria for brain death**

* Fixed pupils which do not respond to sharp changes in the intensity of incident light
* No corneal reflex
* Absent oculo-vestibular reflexes - no eye movements following the slow injection of at least 50ml of ice-cold water into each ear in turn (the caloric test)
* No response to supraorbital pressure
* No cough reflex to bronchial stimulation or gagging response to pharyngeal stimulation
* No observed respiratory effort in response to disconnection of the ventilator for long enough (typically 5 minutes) to ensure elevation of the arterial partial pressure of carbon dioxide to at least 6.0 kPa (6.5 kPa in patients with chronic carbon dioxide retention). Adequate oxygenation is ensured by pre-oxygenation and diffusion oxygenation during the disconnection (so the brain stem respiratory centre is not challenged by the ultimate, anoxic, drive stimulus)

The test should be undertaken by two appropriately experienced doctors on two separate occasions.

1. **Which of the anaesthetic agents below is most likely to induce adrenal suppression?**

|  |  |
| --- | --- |
| A | Propofol |
| B | Etomidate |
| C | Sodium thiopentone |
| D | Ketamine |
| E | Midazolam |

Etomidate is a recognised cause of adrenal suppression, this has been associated with increased mortality when used as a sedation agent in the critically ill.

**Anaesthetic agents**

The table below summarises some of the more commonly used IV induction agents

| **Agent** | **Specific features** |
| --- | --- |
| Propofol | * Rapid onset of anaesthesia * Pain on IV injection * Rapidly metabolised with little accumulation of metabolites * Proven anti emetic properties * Moderate myocardial depression * Widely used especially for maintaining sedation on ITU, total IV anaesthesia and for daycase surgery |
| Sodium thiopentone | * Extremely rapid onset of action making it the agent of choice for rapid sequence of induction * Marked myocardial depression may occur * Metabolites build up quickly * Unsuitable for maintenance infusion * Little analgesic effects |
| Ketamine | * May be used for induction of anaesthesia * Has moderate to strong analgesic properties * Produces little myocardial depression making it a suitable agent for anaesthesia in those who are haemodynamically unstable * May induce state of dissociative anaesthesia resulting in nightmares |
| Etomidate | * Has favorable cardiac safety profile with very little haemodynamic instability * No analgesic properties * Unsuitable for maintaining sedation as prolonged (and even brief) use may result in adrenal suppression * Post operative vomiting is common |

1. **Which measure of central tendency in a dataset is most susceptible to skewness?**

|  |  |
| --- | --- |
| A | Mode |
| B | Median |
| C | Mean |
| D | Both mode and median |
| E | None of the above |

The mean, median and mode are all measures of central tendency. However, the mean is most susceptible to data which is skewed (i.e. has single large outliers). Visual inspection of a dataset (if small) will usually allow a rapid means of determining if this is the case. Plotting data points graphically will also facilitate identification of skewed data. Both median and mode are better alternative measures of central tendency when data is skewed.

**Descriptive statistics**

Descriptive statistics include a point estimate of the measured variable as well as a measure of the variability of the data around that point estimate. Typical examples of point estimates include; mean, median and mode. The two most commonly employed measurements of variability include standard deviation and the inter quartile range. The standard deviation is usually considered in association with the mean, while the inter quartile range is used alongside the median. Other measures of data variability include the standard error of the mean and confidence interval. The standard error of the mean represents the measure of variation around the point estimate of the mean of a group of sample means, as such it should only be used when describing the characteristics of more than one sample.

1. **Which one of the following is equivalent to the pre-test probability?**

|  |  |
| --- | --- |
| A | Post test odds / (1 + post-test odds) |
| B | Pre-test odds x likelihood ratio |
| C | The prevalence of a condition |
| D | The incidence of a condition |
| E | Post-test odds / likelihood ratio |

The prevalence is the proportion of a population that have the condition at a point in time whilst the incidence is the rate at which new cases occur in a population during a specified time period.

**Pre- and post- test odds and probability**

**Pre-test probability**  
The proportion of people with the target disorder in the population at risk at a specific time (point prevalence) or time interval (period prevalence)  
  
For example, the prevalence of rheumatoid arthritis in the UK is 1%  
  
**Post-test probability**  
The proportion of patients with that particular test result who have the target disorder  
  
Post-test probability = post test odds / (1 + post-test odds)  
  
**Pre-test odds**  
The odds that the patient has the target disorder before the test is carried out   
  
Pre-test odds = pre-test probability / (1 - pre-test probability)  
  
**Post-test odds**  
The odds that the patient has the target disorder after the test is carried out  
  
Post-test odds = pre-test odds x likelihood ratio  
  
where the likelihood ratio for a positive test result = sensitivity / (1 - specificity)

1. **A 23 year old man is due to undergo a splenectomy. What is the optimal time for administration of the pneumococcal vaccine?**

|  |  |
| --- | --- |
| A | Two weeks post operatively |
| B | Two weeks pre-operatively |
| C | Six weeks pre-operatively |
| D | Five days pre-operatively |
| E | One month post operatively |

Pre-operative vaccination is preferred, ideally this should be two weeks before surgery.

**Hyposplenism**

Hyposplenism may complicate certain medical conditions where splenic atrophy occurs or may be the result of medical intervention such as splenic artery embolization and splenectomy for trauma. Diagnosis of hyposplenism is difficult and whilst there may be peripheral markers of the splenectomised state (e.g. Howell Jolly bodies) these are neither 100% sensitive or specific. The most sensitive test is a radionucleotide labeled red cell scan.   
Hyposplenism, by whatever mechanism it occurs dramatically increases the risk of post splenectomy sepsis, particularly with encapsulated bacteria. For this reason individuals are recommended to be vaccinated and have antibiotic prophylaxis.   
  
**Key recommendations**

* All those with hyposplenism or may become so (such as prior to an elective splenectomy) should receive pneumococcal, haemophilus type b and meningococcal type C vaccines. These should be administered 2 weeks prior to splenectomy or two weeks following splenectomy. The vaccine schedule for meningococcal disease essentially consists of a dose of Men C and Hib at 2 weeks and then a dose of the MenACWY vaccine one month later. Those aged under 2 may require a booster at 2 years. A dose of pneumococcal polyvalent polysaccharide vaccine (PPV) is given at two weeks. A conjugated vaccine (PCV) is offered to young children. The PCV is more immunogenic but covers fewer serotypes. Boosting PPV is either guided by serological measurements (where available) or by routine boosting doses at 5 yearly intervals.
* Annual influenza vaccination is recommended in all cases
* Antibiotic prophylaxis is offered to all. The risk of post splenectomy sepsis is greatest immediately following splenectomy and in those aged less than 16 years or greater than 50 years. Individuals with a poor response to pneumococcal vaccination are another high risk group. High risk individuals should be counseled to take penicillin or macrolide prophylaxis. Those at low risk may choose to discontinue therapy. All patients should be advised about taking antibiotics early in the case of inter-current infections.
* Asplenic individuals traveling to malaria endemic areas are at high risk and should have both pharmacological and mechanical protection.

**Dosing-antibiotics**  
Penicillin V 500mg BD or amoxicillin 250mg BD  
  
**References**  
Davies J *et al*. Review of guidelines for the prevention and treatment of infection in patients with an absent or dysfunctional spleen: Prepared on behalf of the British Committee for Standards in Haematology by a Working Party of the Haemato-Oncology Task Force. *British Journal of Haematology*2011 (155): 308317.

1. **What is the most appropriate method for describing data generated from an ordinal scale?**

|  |  |
| --- | --- |
| A | Mode and standard deviation |
| B | Mode and inter quartile range |
| C | Mode and standard error of the mean |
| D | Mean and standard deviation |
| E | Median and inter quartile range |

Ordinal data expresses relative differences between subjects when the actual numerical differences are either unknown or cannot be derived. Quantitative comparisons cannot be made for ordinal data. As a result the descriptive statistic of choice for ordinal data is the median and inter quartile range. The inter quartile range describes data between the 25th and 75th centile, with the median illustrating the 50th percentile rank.

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1. **The best graphic representation of frequency distribution data gathered of a continuous variable is:**

|  |  |
| --- | --- |
| A | Line graph |
| B | Simple bar Graph |
| C | Multiple bar chart |
| D | Box Whisker plot |
| E | Histogram |

Histograms are a good method displaying frequencies of continuous variables.

**Continuous variables**

If a variable can take on a value at any point between its minimum and maximum value then it is termed a continuous variable. Otherwise it is called a discrete variable. Patient weights are an example of continuous variables as fractions of numbers are possible (if the equipment is accurate). Discrete variables could include factors such as polyp identification during colonoscopy (as partial polyps don't exist- though partial retrieval certainly can!)

1. **A 53 year old man presents with an ulcerated mass at the anal verge. A biopsy is taken and the histology demonstrates as squamous cell carcinoma. Infection with which of the viruses below is most likely to have contributed to the development of the condition?**

|  |  |
| --- | --- |
| A | Human papillomavirus 16 |
| B | Human T-lymphotropic virus 1 |
| C | Human immunodeficiency virus 2 |
| D | Human immunodeficiency virus 1 |
| E | Human papillomavirus 7 |

Infection with human papilloma virus 16 is a risk factor for the development of intra epithelial dysplasia of the anal skin with subsequent increased risk of invasive malignancy.

**Oncoviruses**

* Viruses which cause cancer
* These may be detected on blood test and prevented by vaccine

These are the main types of oncoviruses and their diseases:

| **Oncovirus** | **Cancer** |
| --- | --- |
| Epstein-Barr virus | Burkitt's lymphoma Hodgkin's lymphoma Post transfusion lymphoma Nasopharyngeal carcinoma |
| Human papillomavirus 16/18 | Cervical cancer Anal cancer Penile cancer Vulval cancer Oropharyneal cancer |
| Human herpes virus 8 | Kaposi's sarcoma |
| Hepatitis B virus | Hepatocellular carcinoma |
| Hepatitis C virus | Hepatocellular carcinoma |
| Human T-lymphotropic virus 1 | Tropical spastic paraparesis Adult T cell leukaemia |

1. **A 54-year-old female is admitted one week following a cholecystectomy with profuse diarrhoea. Apart from a minor intra-operative bile spillage incurred during removal of the gallbladder, the procedure was uncomplicated. What is the most likely diagnosis?**

|  |  |
| --- | --- |
| A | Campylobacter infection |
| B | E. Coli infection |
| C | C. Difficle infection |
| D | Pelvic abscess |
| E | Salmonella infection |

Antibiotics are not routinely administered during an uncomplicated cholecystectomy. Indications for administration of broad spectrum antibiotics include intraoperative bile spillage. Delayed pelvic abscesses following bile spills are extremely rare since most surgeons will manage these intra-operatively.

**Clostridium difficile**

Clostridium difficile is a Gram positive rod often encountered in hospital practice. It produces an exotoxin which causes intestinal damage leading to a syndrome called pseudomembranous colitis. Clostridium difficile develops when the normal gut flora are suppressed by broad-spectrum antibiotics. Clindamycin is historically associated with causing Clostridium difficile but the aetiology has evolved significantly over the past 10 years. Second and third generation cephalosporins are now the leading cause of Clostridium difficile.  
  
Features

* Diarrhoea
* Abdominal pain
* A raised white blood cell count is characteristic
* If severe, toxic megacolon may develop

Diagnosis is made by detecting Clostridium difficile toxin (CDT) in the stool  
  
Management

* First-line therapy is oral metronidazole for 10-14 days
* If severe, or not responding to metronidazole, then oral vancomycin may be used
* For life-threatening infections a combination of oral vancomycin and intravenous metronidazole should be used

1. **In the UK NHS what is the main role of the "Caldicott guardian"?**

|  |  |
| --- | --- |
| A | Liaising between NHS trusts and the General Medical Council about cases of professional misconduct |
| B | Safeguarding the welfare of vulnerable adults |
| C | Investigating cases of suspected child abuse |
| D | Enforcing key principles of the Human Tissue Act |
| E | Safeguarding confidential patient information for an organisation |

**Caldicott guidelines**

A review was commissioned in 1997 by the Chief Medical Officer of England "owing to increasing concern about the ways in which patient information is being used in the NHS in England and Wales and the need to ensure that confidentiality is not undermined. Such concern was largely due to the development of information technology in the service, and its capacity to disseminate information about patients rapidly and extensively".  
  
**Caldicott principles**  
1. **Justify the purpose**  
Every proposed use or transfer of patient identifiable information within or from an organisation should be clearly defined and scrutinised, with continuing uses regularly reviewed, by an appropriate guardian.  
  
2.**Don't use patient identifiable information unless it is absolutely necessary**  
Patient identifiable information items should not be included unless it is essential for the specified purpose(s) of that flow. The need for patients to be identified should be considered at each stage of satisfying the purpose(s).  
  
3.**Use the minimum necessary patient-identifiable information**  
Where use of patient identifiable information is considered to be essential, the inclusion of each individual item of information should be considered and justified so that the minimum amount of identifiable information is transferred or accessible as is necessary for a given function to be carried out.  
  
4.**Access to patient identifiable information should be on a strict need-to-know basis**  
Only those individuals who need access to patient identifiable information should have access to it, and they should only have access to the information items that they need to see. This may mean introducing access controls or splitting information flows where one information flow is used for several purposes.  
  
5.**Everyone with access to patient identifiable information should be aware of their responsibilities**  
Action should be taken to ensure that those handling patient identifiable information - both clinical and non-clinical staff - are made fully aware of their responsibilities and obligations to respect patient confidentiality.  
  
6.**Understand and comply with the law**  
Every use of patient identifiable information must be lawful. Someone in each organisation handling patient information should be responsible for ensuring that the organisation complies with legal requirements.

1. **In order to calculate the sample size for a trial the limits of α and β must be considered in addition to which of the following?**

|  |  |
| --- | --- |
| A | The outcome of interest |
| B | The magnitude and variability of the expected effect size of the intervention on the primary outcome |
| C | The magnitude and variability of the expected effect size of the intervention on the secondary outcome |
| D | All of the above |
| E | None of the above |

In addition to alpha and beta (usually set at 0.05 and 0.1-0.2), investigators need to consider an estimate of the magnitude of and variability of the expected effect size of the intervention on the primary outcome, as well as planned statistical analysis. These data are derived from literature reviews or, where no data is know, from a pilot study.

**Power calculations and statistical error**

**Statistical error**

|  |  |
| --- | --- |
| **Type 1 Error** | A test rejects a true null hypothesis. Analogous to false positive. It usually equates to the significance level assigned to a test. |
| **Type 2 Error** | A test fails to reject a false null hypothesis. It is related to the power of a test. |

**Statistical power**  
The power of a test is the probability that the test will reject the null hypothesis when it is false (thereby avoiding a type 2 error). Increasing the power of a test will reduce the probability of a type 2 error. Usually a value of 0.8 is selected.

1. **A patient attends the clinic having recovered from a major abdominal procedure. At the end of the consultation they hand you £250 GBP in cash. They insist on your taking the gift.   
   What is the best course of action?**

|  |  |
| --- | --- |
| A | Hand the gift to the clinical director |
| B | Accept the gift and document it and the donors details in a formal departmental record |
| C | Hand the gift to the hospital charity |
| D | Pass the gift to a homeless patient on the ward |
| E | Accept the gift and inform the GMC that you have done so |

The GMC guidance on this topic is outlined below.

**Gifts from patients**

The debate surrounding the receipt of gifts is covered by the GMC in their guidance. The following guidance is provided:  
"6. You must not encourage patients to give, lend or bequeath money or gifts that will directly or indirectly benefit you.  
7. You may accept unsolicited gifts from patients or their relatives provided:  
a. this does not affect, or appear to affect, the way you prescribe for, advise, treat, refer, or commission services for patients  
b. you have not used your influence to pressurise or persuade patients or their relatives to offer you gifts.  
8. However, if you receive a gift or bequest from a patient or their relative, you should consider the potential damage this could cause to your patients trust in you and the publics trust in the profession. You should refuse gifts or bequests where they could be perceived as an abuse of trust.  
9. You must not put pressure on patients or their families to make donations to other people or organisations."  
  
They go on to issue the following specific guidance about gifts of monetary value:   
"The acceptance of gifts by general practitioners in all four UK countries is subject to statutory regulation. General Medical Services contract regulations state that a register should be kept of gifts from patients or their relatives which have a value of £100 or more unless the gift is unconnected with the provision of services. The register of gifts should include the donors name and nature of the gift. NHS trusts set their own policies on gifts."

1. A 73 year old man undergoes a brachial embolectomy for an embolus which occurs as a result of atrial fibrillation. He is commence on dabigatran therapy. What is the main mode of action of this drug?

|  |  |
| --- | --- |
| A | Direct inhibition of thrombin |
| B | Activation of antithrombin III |
| C | Inhbition of clotting factors VII and Xa |
| D | Inactivation of clotting factor Xa |
| E | None of the above |

**Anticoagulants**

| **Agent** | **Mechanism of action** | **Monitoring** | **Mechanism of reversal** | **Indications** |
| --- | --- | --- | --- | --- |
| Heparin | Activation of antithrombin III - this inactivates thrombin and factor Xa | APTT | For unfractionated heparin renal inactivation occurs quickly on stopping therapy, for more rapid reversal intravenous protamine may be used | Situations where rapid and predictable anticoagulation is required, low molecular weight heparins are used in stable patients both as prophylaxis against thromboembolic events and also as therapy for these (in higher doses) |
| Warfarin | Inhibits vitamin K dependent synthesis of calcium dependent clotting factors; II, VII, IX, X and protein C and S (the latter two agents are involved in clot degradation and account for the need to maintain heparin therapy whilst treatment is started | INR | Vitamin K, or prothrombin complex concentrate, FFP may also be used but will expand circulating volume | Situations where long term anticoagulation is needed. It is contraindicated in pregnancy |
| Dabigatran | Competitive direct thrombin inhibitor | No monitoring available | Idarucizumab | Prevention of CVA and other embolic events in patients with atrial fibrillation, prevention of DVT in orthopaedic surgery |

1. **A 59 year old women is anxious about having a screening mammogram because of the risk of a false positive examination. Why may such false positives occur?**

|  |  |
| --- | --- |
| A | Test has a high specificity |
| B | Test has a high sensitivity |
| C | There is a high prevalence of breast cancer |
| D | There is a low prevalence of breast cancer |
| E | The test is inaccurate |

**False positive**

A false positive may occur when a screening test falsely identifies individuals as having a condition when none is present. This is a price that is paid for having a sensitive screening test. In surgical practice both faecal occult blood testing and mammography probably generate the greatest burden and worry as a result of false positive results.

1. **A 50 year old man is in a persistent vegetative state following a road traffic accident. He is being cared for in a hospital in England. The consultant is considering withdrawing artificial feeding. What is the most appropriate definitive course of action?**

|  |  |
| --- | --- |
| A | Hold a multidisciplinary meeting and carers and the family and then uphold the consensus decision |
| B | Obtain a second opinion from a consultant in another specialty and then proceed to withdraw if both agree this is the correct course of action |
| C | Approach the English courts for a ruling |
| D | Simply withdraw the feed because there is no clinically perceived benefit and a doctor is not required to continue to provide a treatment in such circumstances |
| E | Withdraw feeding if 3 consultants from unrelated specialties (one of which must be a neurologist) consider withdrawal to be appropriate |

Withdrawal of feeding from patients in PVS in England and Wales is a decision that can only be sanctioned by a court of law.

**Nutrition and end of life decisions**

The General Medical Council has issued guidance about the provision of nutrition in the end of life setting.   
  
The guidance states that the nutritional needs of all patients should be assessed on their own merits. Offer of food and drink by mouth (including spoon feeding) is basic nuture and in UK law is not deemed to be treatment. However, drips, radiological inserted feeding tubes, NG feeding and TPN are deemed to be treatments.  
  
In deciding whether to offer feeding by an invasive means, the patients views should be ascertained. If a patient has capacity and declines such measures then their views must be respected. If the patient does not have capacity, then the risks and benefits of initiating or continuing such treatments should be considered by those caring for the patient.   
  
If a patient requests invasive feeding, but, it is deemed to be clinically inappropriate, then a clinician is not obliged to provide it. However, it is good practice to explore the patients views on the matter carefully and consider obtaining a second opinion where appropriate.  
  
Where a patient is in a persistent vegetative state, the decision to withdraw feeding in England and Wales should be made by a court. The courts in Scotland have not issued such a directive but it is suggested that legal advice be taken in such circumstances. A recent case (Court of Protection, Mr Justice Peter Jackson, M v A Hospital, 20/9/17) has suggested that not all cases need be brought before a court. Particularly when the views of health care professionals and family are aligned. However, this is not yet, published GMC guidance. Where doubt exists, a court application remains likely.

1. **Theme:** Body mass index

|  |  |
| --- | --- |
| **A.** | Obese Class III (Very severely obese) |
| **B.** | Very severely underweight |
| **C.** | Normal |
| **D.** | Obese Class I (Moderately obese) |
| **E.** | Severely underweight |
| **F.** | Underweight |
| **G.** | Overweight |
| **H.** | Obese Class II (Severely obese) |

Please match the following body mass indexes to the descriptors provided. Each option may be used once, more than once or not at all.

|  |  |
| --- | --- |
| **1.** | BMI 20  http://d35pzpz673ryzz.cloudfront.net/css/small_tick.gif Normal |
| **2.** | BMI 24  http://d35pzpz673ryzz.cloudfront.net/css/small_tick.gif Normal |
| **3.** | BMI 37  http://d35pzpz673ryzz.cloudfront.net/css/small_tick.gif Obese Class II (Severely obese) |

**Body mass index**

Body mass index is widely used and is calculated by the formula;  
BMI= mass (Kg)/height (m)2  
  
**Values**

| **Category** | **Value** |
| --- | --- |
| Very severely underweight | less than 15 |
| Severely underweight | from 15.0 to 16.0 |
| Underweight | from 16.0 to 18.5 |
| Normal (healthy weight) | from 18.5 to 25 |
| Overweight | from 25 to 30 |
| Obese Class I (Moderately obese) | from 30 to 35 |
| Obese Class II (Severely obese) | from 35 to 40 |
| Obese Class III (Very severely obese) | over 40 |

Obesity classification is taken from the WHO obesity classification system.

1. **Which of the following statements relating to MRI scanning is untrue?**

|  |  |
| --- | --- |
| A | They apply a field strength that is measured in teslas |
| B | Co-administration of gadolinium is usually required for accurate imaging of vessels |
| C | In T1 images fat is darker than water |
| D | It is safe to use in pregnancy |
| E | Gadolinium is usually used to enhance T1 images |

T1 images water is dark and fat bright  
T2 images fat is dark and water bright

**MRI scanning**

* Non radioactive imaging technique involving application of electromagnetic field to tissues resulting in proton spinning
* Magnetic field has uniform field density and strength
* Field strength measured in teslas,open magnets have lower teslas
* Gadolinium based contrast agents may improve resolution
* Most MRI contrast agents work by reducing the T1 relaxation times of neighboring protons with an increased rate of stimulated emission from high energy states
* Gadolinium has a better safety profile than iodonated contrast media
* Basic scans are referred to as either T1 or T2 weighted and this refers to the spin lattice relaxation time
* T1 scans are often collected before and after infusion of contrast agents and are useful for distinguishing between grey and white matter. In T1 images water is dark and fat is bright.
* In T2 scans, the water fat density is reversed so that water is bright and fat is dark
* It is safe to use in pregnancy. Implanted devices are contra indications to use

1. **Which of the following is least commonly associated with Dupuytrens contracture?**

|  |  |
| --- | --- |
| A | Peyronie's disease |
| B | Alcoholic cirrhosis |
| C | Phenytoin use |
| D | Mycobacterium tuberculosis infection |
| E | Dysplasia of the palmar fascia |

In Dupuytrens disease the palmar fascia becomes hyperplastic and subsequently contracts. It is associated with the condition Peyronies disease in which the penis may become distorted. It is associated with liver disease, drugs such as phenytoin which can induce epithelial hyperplasia and chronic infections. A number of surgical excisional therapies are described and should be reserved for those with progressive or debilitating symptoms.

*Dupuytrens contracture*

* Fixed flexion contracture of the hand where the fingers bend towards the palm and cannot be fully extended.
* Caused by underlying contractures of the palmar aponeurosis . The ring finger and little finger are the fingers most commonly affected. The middle finger may be affected in advanced cases, but the index finger and the thumb are nearly always spared.
* Progresses slowly and is usually painless. In patients with this condition, the tissues under the skin on the palm of the hand thicken and shorten so that the tendons connected to the fingers cannot move freely. The palmar aponeurosis becomes hyperplastic and undergoes contracture.
* Commonest in males over 40 years of age.
* Association with liver cirrhosis and alcoholism (there is a historical association with TB infection). However, many cases are idiopathic.
* Treatment is surgical and involves fasciectomy. However, the condition may recur and many surgical therapies are associated with risk of neurovascular damage to the digital nerves and arteries.

1. **What is the mechanism of action of ciprofloxacin?**

|  |  |
| --- | --- |
| A | Destruction of bacterial aquaporin proteins |
| B | Inhibition of reverse transcriptase |
| C | Direct injury to the bacterial cell wall |
| D | Inhibition of DNA gyrase |
| E | Osmotic injury to the cell |

**Antibiotics : mechanism of action**

The lists below summarise the site of action of the commonly used antibiotics  
  
Inhibit cell wall formation

* penicillins
* cephalosporins

Inhibit protein synthesis

* aminoglycosides (cause misreading of mRNA)
* chloramphenicol
* macrolides (e.g. erythromycin)
* tetracyclines
* fusidic acid

Inhibit DNA synthesis

* quinolones (e.g. ciprofloxacin)
* metronidazole
* sulphonamides
* trimethoprim

Inhibit RNA synthesis

* rifampicin

1. **Theme:** Sutures

|  |  |
| --- | --- |
| **A.** | 3/0 undyed polybutester |
| **B.** | 3/0 undyed polyglactin |
| **C.** | 1/0 dyed polyglyconate |
| **D.** | 1 dyed polydioxanone |
| **E.** | 5/0 polypropylene |
| **F.** | 3/0 dyed polyglyconate |
| **G.** | 4/0 silk |
| **H.** | 6/0 polyester |
| **I.** | 2/0 undyed polybutester |
| **J.** | 1 dyed polyglycolic acid |

Please select the most appropriate suture for the situation described. Each option may be used once, more than once or not at all.

|  |  |
| --- | --- |
| **1.** | Closing the linea alba  http://d35pzpz673ryzz.cloudfront.net/css/small_tick.gif 1 dyed polydioxanone  1 PDS is commonly used for this purpose as it has the tensile strength. Note that polyglycolic acid (Dexon) will not retain its tensile strength for long enough. |
| **2.** | Construction of an ileo-colic anstomosis  **The correct answer is 3/0 dyed polyglyconate**  A hand sewn ileo-colic anastomosis will typically use a 3/0 suture as this will provide adequate tensile strength. Many sutures could be used but only polyglyconate has the correct size of the options provided. |
| **3.** | Placing a patch onto the femoral artery  **The correct answer is 5/0 polypropylene**  Polypropylene is a favored for vascular suturing because it is inert and retains its tensile strength. Polyester also fulfills this requirement. However, the suture sizes in the options do not match. |

**Sutures**

The ideal sutures should fulfill the following criteria:

* Achieve its purpose
* Disappear once its work is complete
* Be free from infection
* Be non- irritant

(Moynihan 1920)  
  
Many sutures are currently available and for the purposes of examinations it is important to be clear and distinct on the difference between trade names and raw material/ generic names. Commonly used sutures are described in the table below;

| **Agent** | **Classification** | **Durability** | **Uses** | **Special points** |
| --- | --- | --- | --- | --- |
| Silk | Braided  Biological | Theoretically permanent although strength not preserved | Anchoring devices, skin closure | Knots easily, poor cosmesis |
| Polyglactin 910 (vicryl) | Braided multifilament, synthetic | Strength retained for 60% at 2 weeks, broken down by hydrolysis , complete by 90 days | Widespread, ranging from visceral anastomosis through to skin closure | Use undyed for skin closure |
| Polyglyconate (maxon) | Synthetic monofilament | Tensile strength 70% at 2 weeks, complete absorption by 180 days | Used as alternative to PDS and polyglactin 910 in some centres | Undyed version available |
| Polydiaxanone (PDS) | Synthetic , polyester polymer | 70% tensile strength at 2 weeks, breakdown up to 3 months (longer with thicker sutures) | Widespread surgical applications including visceral anastomoses, dermal closure, mass closure of abdominal wall | Used in most surgical specialties (avoid dyed form in dermal closure) |
| Polyester (ethibond, Dacron, tecron) | Both monofilament and braided synthetic | Remains indefinitely, tensile strength preserved | Vascular grafts, prolonged tissue approximation | n/a |
| Polypropylene (prolene) | Synthetic monofilament | Both suture and tensile strength persist indefinitely | Especially popular for vascular anastomoses | Poor handling characteristics |
| Nylon (ethilon) | Synthetic mono and multifilament | Degrades at a rate of 15% per year | Wide range of uses | Avoid in vascular anastomoses |

When catgut was withdrawn from use Ethicon introduced vicryl rapide sutures, these have a polymer of lower molecular weight that is more rapidly hydrolysed and therefore absorbs more quickly.  
  
**Sizes**

| **USP gauge** | **Diameter (max in mm** |
| --- | --- |
| 10/0 | 0.025 |
| 4/0 | 0.203 |
| 2/0 | 0.330 |
| 0 | 0.406 |
| 1 | 0.483 |
| 2 | 0.559 |

1. **Which of the following blood products can be administered to a non ABO matched recipient?**

|  |  |
| --- | --- |
| A | Whole blood |
| B | Platelets |
| C | Packed red cells |
| D | Fresh frozen plasma |
| E | Cryoprecipitate |

In the UK, platelets either come from pooling of the platelet component from four units of whole donated blood, called random donor platelets, or by plasmapharesis from a single donor. The platelets are suspended in 200-300 ml of plasma and may be stored for up to 4 days in the transfusion laboratory where they are continually agitated at 22oC to preserve function. One adult platelet pool raises the normal platelet count (150 - 450 platelets x 109/litre) by 510 platelets x 109/litre. ABO identical or compatible platelets are preferred but not necessary in adults; but rhesus compatibility is required in recipients who are children and women of childbearing age to prevent haemolytic disease of the newborn.

**Cross matching blood products**

**Cross matching**

| **Must be cross matched** | **Can be ABO incompatible in adults** |
| --- | --- |
| Packed red cells | Platelets |
| Fresh frozen plasma |  |
| Cryoprecipitate |  |
| Whole blood |  |

1. **A 45-year-old man presents to surgical outpatients with a long history of recurrent abdominal pain and vomiting. He is noted to have a peripheral motor neuropathy on examination. What is the most likely diagnosis?**

|  |  |
| --- | --- |
| A | Huntington's disease |
| B | Myeloma |
| C | Acute intermittent porphyria |
| D | Lawrence-Moon-Biedl syndrome |
| E | Friedreich's ataxia |

Neurological signs combined with abdominal pain is acute intermittent porphyria or lead poisoning until proven otherwise.  
Lawrence-Moon-Biedl syndrome is a pleiotropic disorder with variable expressivity and a wide range of clinical variability observed both within and between families. The main clinical features are rodcone dystrophy, with childhood-onset visual loss preceded by night blindness; postaxial polydactyly; truncal obesity that manifests during infancy and remains problematic throughout adulthood; specific learning difficulties in some but not all individuals; male hypogenitalism and complex female genitourinary malformations; and renal dysfunction, a major cause of morbidity and mortality.

**Acute intermittent porphyria**

Acute intermittent porphyria (AIP) is a rare autosomal dominant condition caused by a defect in porphobilinogen deaminase, an enzyme involved in the biosynthesis of haem. The results in the toxic accumulation of delta aminolaevulinic acid and porphobilinogen. It characteristically presents with abdominal and neuropsychiatric symptoms in 20-40 year olds. AIP is more common in females (5:1)  
  
**Features**

* abdominal: abdominal pain, vomiting
* neurological: motor neuropathy
* psychiatric: e.g. depression
* hypertension and tachycardia common

**Diagnosis**

* classically urine turns deep red on standing
* raised urinary porphobilinogen (elevated between attacks and to a greater extent during acute attacks)
* assay of red cells for porphobilinogen deaminase
* raised serum levels of delta aminolaevulinic acid and porphobilinogen

1. **Which of the following is not utilised as a descriptive statistic?**

|  |  |
| --- | --- |
| A | Mean |
| B | Median |
| C | Mode |
| D | Z score |
| E | Standard deviation |

The z score is determined using the normal distribution and is not a descriptive statistic.

**Descriptive statistics**

Descriptive statistics include a point estimate of the measured variable as well as a measure of the variability of the data around that point estimate. Typical examples of point estimates include; mean, median and mode. The two most commonly employed measurements of variability include standard deviation and the inter quartile range. The standard deviation is usually considered in association with the mean, while the inter quartile range is used alongside the median. Other measures of data variability include the standard error of the mean and confidence interval. The standard error of the mean represents the measure of variation around the point estimate of the mean of a group of sample means, as such it should only be used when describing the characteristics of more than one sample.

1. **Which of the following is most useful in distinguishing between liver metastases from colorectal cancer and small liver haemangioma on CT scanning?**

|  |  |
| --- | --- |
| A | Hypovascular |
| B | Multifocal nature |
| C | Size |
| D | Hypodense relative to surrounding liver tissue |
| E | None of the above |

Haemangioma remains an important differential diagnosis and confirmation with MRI (which accurately identifies such lesions) should occur prior to surgery.

**Liver metastasis from colorectal cancer**

Approximately 70% of patients with metastatic colorectal cancer will have disease that is confined to the liver.   
Detection is usually made using CT scanning. Colorectal metastases will usually be hypovascular relative to the surrounding liver tissue and appear to be hypoattentuating on CT. On MRI scanning they will usually appear as hypodense lesions on T1 weighted image and hyperdense on T2 weighted images. Only 15% of patients will have disease that is surgically resectable.   
  
**Classification of resectable disease**

| **Resection category** | **Features** |
| --- | --- |
| Usually resectable | Four or fewer segments or deposits in the liver Residual liver volume >40% Vena cava not involved Contra lateral portal pedicle |
| Potential resection | Involvement of 5-6 segments Contra lateral named vascular structure involvement Central hepatectomy Vascular reconstruction |
| Not resectable | Involvement of two portal branches Involvement of three hepatic veins Marked extra hepatic disease (e.g. portal nodes, non resectable distant disease) |

**Role of chemotherapy**  
Use of FOLFOX 4 chemotherapy regime is standard. The agents used include; oxaliplatin, fluorouracil and folinic acid. This is typically given prior to liver resection. A regime lasting 3 months is usually favored as it provides the best compromise between treatment related toxicity and improvement in outcome.   
Recurrence is seen in up to 60% of patients undergoing surgical resection of liver metastasis. Usually within the first 1-2 years.

1. **Which of the following statements relating to transfusion related lung injury is untrue?**

|  |  |
| --- | --- |
| A | It is commoner following transfusion with fresh frozen plasma than with packed red cells |
| B | It is a recognised complication of platelet transfusion |
| C | It occurs as a result of leucocyte antibodies in the transfused plasma |
| D | Typically manifests as acute onset pulmonary oedema 1-2 hours following transfusion |
| E | Early therapy with diuretics reduces pulmonary infiltrates and improves outcomes |

Transfusion related lung injury is most common following the transfusion of plasma components such as platelets and FFP. The condition is due to leucocyte antibodies in the transfused plasma. This causes leucocyte sequestration and degranulation in the lung. This produces marked microvascular and tissue damage with the development of a non cardiogenic pulmonary oedema. Because the primary problem is one of tissue injury, diuretic therapy is largely unhelpful.

**Massive haemorrhage**

**Definition**  
This is the loss of one blood volume in a 24 hour period or the loss of 50% of the circulating blood volume in 3 hours. A blood loss of 150ml/ minute is also included. The normal adult blood volume is 7% of total adult body weight. The blood volume equates to between 8 and 9% of a child's body weight.   
  
**Complications of massive transfusion**

| **Complication** | **Key points** |
| --- | --- |
| Hypothermia | Blood is refrigerated Hypothermic blood impairs homeostasis  Shifts Bohr curve to the left |
| Hypocalcaemia | Both FFP and platelets contain citrate anticoagulant, this may chelate calcium |
| Hyperkalaemia | Plasma of red cells stored for 4-5 weeks contains 5-10 mmol K+ |
| Delayed type transfusion reactions | Due to minor incompatibility issues especially if urgent or non cross matched blood used |
| Transfusion related lung injury | Acute onset non cardiogenic pulmonary oedema Leading cause of transfusion related deaths Greatest risk posed with plasma components Occurs as a result of leucocyte antibodies in transfused plasma Aggregation and degranulation of leucocytes in lung tissue accounts for lung injury |
| Coagulopathy | Anticipate once circulating blood volume transfused 1 blood volume usually drops platelet count to 100 or less 1 blood volume will both dilute and not replace clotting factors Fibrinogen concentration halves per 0.75 blood volume transfused |

**References**  
Stainsby *et al*. Guidelines on the management of massive blood loss. *British Journal of Haematology*2006 (135): 534 641.