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01305 361132 or via dhc.infectionprevention.control@nhs.net. This is an alternative to a PDF document to assist with user accessibility. This is an evidence-based practice manual for use by all those involved in care provision in England. It should be adopted as mandatory guidance in NHS settings or settings where NHS services are delivered, and the principles should be applied in all care settings. The manual ensures consistent UK-wide approach to infection prevention and control, although some operational and organisational details may differ across the nations. Accessed version 2.1.1 Updated 24 April 2025. Standard infection prevention and control precautions (SICPs) are to be used by all staff, in all care settings, at all times, for all patients whether infection is known to be present or not, to ensure the safety of those being cared for, staff and visitors in the care environment. SICPs are the basic infection prevention and control measures necessary to reduce the risk of transmitting infectious agents from both recognised and unrecognised sources of infection. Sources of (potential) infection include blood and other body fluids, secretions or excretions (excluding sweat), non-intact skin or mucous membranes and any equipment or items in the care environment that could have become contaminated. The application of SICPs during care delivery is determined by assessing risk to and from individuals. This includes the task, level of interaction and/or the anticipated level of exposure to blood and/or other body fluids. To protect effectively against infection risks, SICPs must be used consistently by all staff. SICPs implementation monitoring must also be ongoing to ensure compliance with safe practices and to demonstrate ongoing commitment to patient, staff and visitor safety as required by the Health and Safety Executive and the care regulators, the Care Quality Commission. There are 10 elements of SICPs: patient placement/assessment of infection risk hand hygiene respiratory and cough hygiene personal protective equipment safe management of the care environment safe management of care equipment safe management of healthcare linen safe management of blood and body fluids safe disposal of waste (including sharps) occupational safety/managing prevention of exposure (including sharps) 1.1 Patient placement/assessment for infection risk Patients must be promptly assessed for infection risk on arrival at the care area, eg inpatient/outpatient/care home, (if possible, prior to accepting a patient from another care area) and should be continuously reviewed throughout their stay. This assessment should influence placement decisions in accordance with clinical/care needs(s). Patients who may present a cross-infection risk include those with diarrhoea, vomiting, an unexplained rash, fever or respiratory symptoms (which have been previously positive with a multi-drug resistant organism (MDRO), eg MRSA, CPE who have been an inpatient in any hospital in the UK or abroad or are a known epidemiological link to a carrier of CPE. The National Infection Prevention and Control Isolation Prioritisation Tool provides a systematic framework that can be used to assist in the prioritisation of isolation rooms as part of multidisciplinary assessment. NB the Isolation Prioritisation tool MUST not be used for patients with a suspected High Consequence Infectious Disease (HCID). Further information can be found in the patient placement literature review. 1.2 Hand hygiene Hand hygiene is considered one of the most important ways to reduce the transmission of infectious agents that cause healthcare associated infections (HCIs). Clinical hand-wash basins must: be used for that purpose only and not used for the disposal of other liquids had mixer taps, no overflow or plug and be in a good state of repair have wall mounted liquid soap and paper towel dispensers. Hand hygiene facilities should include instructional posters. Before performing hand hygiene: expose forearms (bare below the elbow). If disposable over-sleeves are worn for religious reasons, these must be removed and disposed of before performing hand hygiene, then replaced with a new pair\* removed all hand and wrist jewellery. The wearing of a single, plain metal finger ring, eg a wedding band, is permitted but should be removed (or moved up) during hand hygiene. A religious bangle can be worn but should be moved up the forearm during hand hygiene and secured during patient care activities ensure fingernails are clean and short, and do not wear artificial nails or nail products cover all cuts or abrasions with a waterproof dressing. \*refer to NHS England uniforms and workwear guidance (Appendix B) for more information on the use of over-sleeves and longer-sleeved uniforms. To perform hand hygiene: Wash hands with non-antimicrobial liquid soap and water if: hands are visibly soiled or dirty caring for patients with vomiting or diarrhoeal illnesses caring for a patient with a suspected or known gastrointestinal infection, eg norovirus or a spore-forming organism such as clostridioides difficile. In all other circumstances, use alcohol-based handrubs (ABHRs) for routine hand hygiene during care. ABHRs must be available for staff as near to the point of care as possible. Where this is not practical, personal ABHR dispensers should be used, eg within the community, domiciliary care, mental health units etc. In settings where personal ABHR dispensers are deemed unsuitable due to staff safety concerns, organisations consider alternative products and are responsible for ensuring safe systems of work, including the completion of a documented risk assessment approved through local governance procedures. Organisations must confirm the efficacy and suitability of the product (i.e., that it conforms with the relevant standards and is appropriate for the intended use) with the product manufacturer. Any differences in use and application, including volume, contact and disinfection time, of an alternative product compared with ABHR should be identified as part of this assessment and appropriate implementation plans should include education and supporting materials for staff. Where running water is unavailable, or hand hygiene facilities are lacking, staff may use hand wipes followed by ABHR and should wash their hands at the first opportunity. Perform hand hygiene: before touching a patient, before clean or aseptic procedures, after body fluid exposure risk after touching a patient; and after touching a patient's immediate surroundings. Always perform hand hygiene before putting on and after removing gloves. For how to wash hands, see the step-by-step guide in appendix 1 of this document. For how to hand rub, see the step-by-step guide in appendix 2 of this document. Skin care dry hands thoroughly after hand washing, using disposable paper towels use an emollient hand cream regularly eg during breaks and when off duty do not use or provide communal tubs of hand cream in the care setting staff with skin problems should seek advice from occupational health or their GP and depending on their skin condition and the severity may require additional interventions or reporting. Surgical hand antiseptis Surgical scrubbing/rubbing (this applies to those undertaking surgical and some invasive procedures): perform surgical scrubbing/rubbing before donning sterile theatre garments or at other times, eg before inserting central vascular access devices remove all hand and wrist jewellery (including wedding band) nail brushes should not be used for surgical hand antiseptis nail picks (single-use) can be used if nails are visibly dirty soft, non-abrasive, sterile (single-use) sponges may be used to apply antimicrobial liquid soap to the skin if licensed for this purpose use an antimicrobial liquid soap licensed for surgical scrubbing or an ABHR licensed for surgical rubbing (as specified on the product label) ABHR can be used between surgical procedures if licensed for this use or between glove changes if hands are not visibly soiled. For surgical scrubbing (not rubbing), follow the step-by-step guide in appendix 3 of this document. For surgical rubbing (not scrubbing), follow the step-by-step guide in appendix 4 of this document. For hand hygiene posters/leaflet, refer to the resources section of NIPCM. Further information in the hand hygiene literature reviews: 1.3 Respiratory and cough hygiene Respiratory and cough hygiene is designed to minimise the risk of cross transmission of known or suspected respiratory illness (pathogens): cover the nose and mouth with a disposable tissue when sneezing, coughing, wiping and blowing the nose; if unavailable use the crook of the arm disposed of all used tissues promptly into a waste bin wash hands with non-antimicrobial liquid soap and warm water after coughing, sneezing, using tissues, or after contact with respiratory secretions or objects contaminated by these secretions where there is no running water available or hand hygiene facilities are lacking, staff may use hand wipes followed by ABHR and should wash their hands at the first available opportunity keep contaminated hands away from the eyes nose and mouth. Staff should promote respiratory and cough hygiene helping those (eg, elderly, children) who need assistance with this, eg providing patients with tissues, a dedicated receptacle i.e. waste bag for used tissues and hand hygiene facilities as necessary. Further information can be found in cough etiquette/respiratory hygiene in the hospital setting literature review. 1.4 Personal protective equipment (PPE) Before undertaking any procedure, staff should assess any likely exposure to blood and/or other body fluids, non-intact skin or mucous membranes and wear personal protective equipment (PPE) that protects adequately against the risks associated with the procedure. The principles of PPE use set out below are important to ensure that PPE is used correctly to ensure patient and staff safety. Avoiding overuse or inappropriate use of PPE is a key principle that ensures this is risk-based and minimises its environmental impact. Where appropriate, consideration should be given to the environmental impact of sustainable or reusable PPE options versus single-use PPE while adhering to the principles below. All PPE must be: located close to the point of use, PPE for healthcare professionals providing care in the community and domiciliary care providers must be transported in a clean receptacle stored to prevent contamination in a clean, dry area until required (expiry dates must be adhered to) single-use only unless specified by the manufacturer changed immediately after each patient and/or after completing a procedure or task disposed of after use into the correct waste stream, eg domestic waste, offensive (non-infectious) or clinical waste discarded if damaged or contaminated. NB Reusable PPE such as goggles/face shields/visors, must be decontaminated after each use according to manufacturer's instruction. Gloves must be: worn when exposure to blood and/or other body fluids, non-intact skin or mucous membranes is anticipated or likely changed immediately after each patient and/or after completing a procedure/task even on the same patient, and hand hygiene performed changed if a perforation or puncture is suspected appropriate for use, fit for purpose and well-fitting never decontaminated with ABHR or soap between use low risk of causing sensitisation to the wearer appropriate for the tasks being undertaken, taking into account the substances being handled, type and duration of contact, size and comfort of the gloves, and the task and requirement for glove robustness and sensitivity. Sterile gloves must be worn: when sterility is required in an operating theatre, and for some aseptic techniques eg insertion of central venous catheters, insertion of peripherally inserted central catheters, insertion of pulmonary artery catheters and spinal, epidural and caudal procedures NB Double gloving is NOT recommended for routine clinical care. However, it may be required for some exposure prone procedures, eg orthopaedic and gynaecological operations, when attending major trauma incidents or as part of additional precautions for high consequence infectious disease management. Gloves are NOT required to carry out near patient administrative tasks, eg when using the telephone, using a computer or tablet, writing in the patient chart, eg when performing surgical procedures or epidurals or inserting a central vascular catheter (CVC) (Type II (not classed as an FRSM) or Type IIR) well-fitting and fit for purpose, fully covering the mouth and nose (manufacturers' instructions must be followed to ensure effective fit and protection) removed or changed – at the end of a procedure/task contaminated or soiled. Aprons must be: worn to protect uniform or clothes when contamination is anticipated or likely changed between patients and/or after completing a procedure or task. Full body gowns or fluid-resistant coveralls must be: worn when there is a risk of extensive splashing of blood and/or body fluids, eg operating theatre, ITU worn when a disposable apron provides inadequate cover for the procedure or task being performed changed between patients and removed immediately after completing a procedure or task sterile when sterility is required in an operating theatre and for some aseptic techniques eg for insertion of central venous catheters, insertion of peripherally inserted central catheters, insertion of pulmonary artery catheters and spinal, epidural and caudal procedures. Further information can be found in the aprons/gowns literature review. Eye or face protection (including full-face visors) must: be worn if blood and/or body fluid contamination to the eyes or face is anticipated or likely, eg by members of the surgical theatre team and always during aerosol generating procedures; regular corrective spectacles are not considered eye protection not be impeded by accessories such as piercings or false eyelashes not be touched when being worn. Further information can be found in the eye/face protection literature review. Fluid resistant surgical face masks (FRSM): Surgical face masks are required: as a means of source control, eg to protect the patient from the wearer during sterile procedures such as surgery, and to protect the wearer when there is a risk splashing or spraying of blood, body fluids, secretions or excretions onto the respiratory mucosa, as an element of PPE for droplet precautions (see section 2.4 and appendices 5b and c). FRSM must be worn (with eye protection) if a full-face visor is not available and spraying or splashing of blood, body fluids, secretions or excretions onto the respiratory mucosa (nose and mouth) is anticipated or likely (Type IIR) worn to protect patients from the operator as a source of infection, eg when performing surgical procedures or epidurals or inserting a central vascular catheter (CVC) (Type II (not classed as an FRSM) or Type IIR) well-fitting and fit for purpose, fully covering the mouth and nose (manufacturers' instructions must be followed to ensure effective fit and protection) removed or changed – at the end of a procedure/task – if the mask's integrity is breached, eg from moisture build-up after extended use or from gross contamination with blood or body fluids – in accordance with manufacturers' specific instructions. Further information can be found in the surgical face masks literature review. Footwear must be: visibly clean, non-slip and well-maintained, and support and cover the entire foot to avoid contamination with blood or other body fluids or potential injury from sharps removed before leaving a care area where dedicated footwear is used, eg theatre; these areas must have a decontamination schedule with responsibility assigned. Further information can be found in the footwear literature review. Headwear Headwear is not routinely required in clinical areas unless part of theatre attire or to prevent contamination of the environment such as in clean rooms. Headwear must be: worn in theatre settings and clean rooms, eg central decontamination unit well-fitting and completely cover the hair changed or disposed of between clinical procedures/lists or tasks and if contaminated with blood and/or body fluids removed before leaving the theatre or clean room individuals with facial hair must also cover this in areas where headwear is required, eg wear a snood. NB Headwear worn for religious reasons such as turbans, kippot veils, headscarves must not compromise patient care and safety. These must be washed and/or changed daily or immediately if contaminated and comply with additional attire requirements, for example, in theatres. Further information can be found in the headwear literature review. For the recommended method of putting on and removing PPE, refer to appendix 6. 1.5 Safe management of care equipment Care equipment is easily contaminated with blood, other body fluids, secretions and infectious agents. Consequently, it is easy to transfer infectious agents from communal care equipment during care delivery. Care equipment is classified as either: single-use: equipment which is used once on a single patient then discarded. This equipment must never be re-used. The packaging will carry the symbol of the number two in a circle with a diagonal cross single patient use: equipment which can be reused on the same patient and may require decontamination in-between use such as nebuliser, mask reusable invasive equipment: used once then decontaminated, eg surgical instruments and solid state reusable equipment, such as, flexible endoscopes and transducers reusable non-invasive equipment: (often referred to as communal equipment) – reused on more than one patient following decontamination between each use, eg commode, patient transfer trolley. NB Needles and syringes are single use devices, they should never be used more than once or reused to draw up additional medication. Never administer medications from a single-dose vial or intravenous (IV) bag to multiple patients. Before using any sterile equipment check that: the packaging is intact there are no obvious signs of packaging contamination the expiry date remains valid any sterility indicators are consistent with the process being completed successfully. Decontamination of reusable non-invasive care equipment must be undertaken: between each use/between patients after blood and/or body fluid contamination at regular predefined intervals as part of an equipment cleaning protocol before inspection, servicing or repair. If providing domiciliary care, equipment should be transported safely and decontaminated as above before leaving the patient's home. Always adhere to Control of Substances Hazardous to Health (COSHH) risk assessments and manufacturers' guidance for use and decontamination of all care equipment. All reusable non-invasive care equipment must be decontaminated between patients/clients using either approved detergent wipes or detergent solution, in line with manufacturers' instructions, before being stored clean and dry. Decontamination protocols must include responsibility for; frequency of; and method of environmental decontamination an equipment decontamination status certificate will be required if any item of equipment is being sent to a third party, eg for inspection, servicing or repair guidance should be sought from the infection prevention and control team prior to procuring, trialling or sending any reusable non-invasive equipment medical devices and other care equipment must have evidence of planned preventative maintenance programmes. For how to decontaminate reusable non-invasive care equipment see Appendix 7. For decontamination of surgical instruments see HTM01-01 decontamination of surgical instruments. Further information can be found in the management of patient care equipment literature review. 1.6 Safe management of the care environment The care environment must be: Always adhere to COSHH risk assessments for product use and processes for decontamination of the care environment. Routine cleaning the environment should be routinely cleaned in accordance with the National Cleaning Standards use of detergent wipes is acceptable for cleaning surfaces/frequently touched sites within the care area a fresh solution of general-purpose neutral detergent in warm water is recommended for routine cleaning. This should be changed when dirty or when changing tasks routine disinfection of the environment is not recommended however, 1,000ppm available chlorine should be used routinely on sanitary fittings/staff groups should be aware of their environmental cleaning schedules for their area and clear on their specific responsibilities cleaning protocols should include responsibility for, frequency of, and method of environmental decontamination. Further information can be found in the safe management of the care environment literature review. 1.7 Safe management of linen Healthcare laundry must be managed and segregated in accordance with HTM 01-04 . Healthcare linen is categorised as: Clean linen - linen washed and ready to be used. Used (soiled and fouled) linen - used linen, irrespective of state, which on occasion may be contaminated by blood or body fluids, and Infectious linen - linen that has been used by a patient who is known or suspected to be infectious. Storage and handling of clean linen: Hand hygiene should be performed prior to handling clean linen. Clean linen should be removed from plastic bags before storage to prevent the growth of Bacillus cereus. Clean linen should be stored above floor level in a designated area, preferably an enclosed or cupboard that is clean, dry and cool. If clean linen is not stored in a cupboard, then the trolley used for storage must be designated for this purpose and completely covered with an impervious covering/door that is able to withstand decontamination. Clean linen storage areas should be dedicated for the purpose and appropriately designed to prevent damage to linen and to allow for the rotation of stocks. Handling linen should be physically separated from used/infectious linen when in storage and during transport. Storage and handling of used (previously known as soiled/fouled linen) and infectious linen: Staff handling used and/or infectious linen must wear appropriate PPE (see section 1.4). Hand hygiene must be performed after handling used and/or infectious linen. Ensure a laundry receptacle is available as close as possible to the point of use for immediate linen deposit. Used items of linen should be removed one by one and placed in the used linen hamper/stream. Do not: rinse, shake or sort linen on removal from beds/trolleys placed used linen on the floor or any other surfaces eg a locker/table top re-handle used linen once bagged overflow laundry receptacles (not more than 2/3 full); or place inappropriate items in the laundry receptacle eg used equipment/needles Infectious linen must not be sorted but should be rolled together and sealed in a water-soluble bag (entirely water soluble 'alginate' bag or impermeable bag with soluble seals), which is then placed in an impermeable bag immediately on removal from the bed and secured before leaving a clinical area. Linen should be placed in an impermeable bag immediately on removal from the bed or before leaving a clinical department Linen bags/receptacles must be tagged (eg, hospital ward/care area) and dated Store all used/infectious linen in a designated, safe, lockable area while awaiting collection. Collection schedules must be acceptable to the care area and there should be no build-up of linen receptacles All linen that is deemed unfit for re-use, eg, torn or heavily contaminated, should be categorised at the point of use and returned to the laundry for assessment and disposal. Linen used during patient transfer, eg, blankets, should be categorised at the point of destination. Linen from patients infected with, or at high risk of having, Hazard Group 4 organisms (haemorrhagic fever viruses such as Lassa Fever) should be placed in a Category A waste and must not be returned to a laundry. Further information can be found in the safe management of linen literature review. For how to manage linen at care area level see Appendix 8. 1.8 Safe management of blood and body fluid spillages Spillages of blood and other body fluids may transmit blood borne viruses. Spillages must be treated immediately by staff trained to undertake this safely. Responsibilities for the management of blood/body fluid spills must be clear within each area/care setting. For management of blood and body fluid spillages see Appendix 9. If an organisation locally approves a product for use in the management of blood and body fluid spills, the organisation is responsible for ensuring safe systems of work, including the completion of a documented risk assessment approved through local governance procedures. Organisations must confirm the efficacy and suitability of the product (i.e., that it conforms with the relevant standards and is appropriate for the intended use) with the product manufacturer. A locally approved product which conforms to: EN17126, EN13727, EN14348, EN14476, EN14885, EN13697, EN14885, EN13704, EN1650, EN1276 and EN13624 may be used for the management of blood and body fluid spills. Further information can be found in the management of blood and body fluid literature review. Healthcare providers should ensure that any polymer gel for non-patient use (eg spill kits, controlled drug destruction, use by cleaning staff) is kept secure and away from patients. See National Patient Safety Alert - National Patient Safety Alert - Superabsorbent polymer gel granules (2019) NatPSA/2019/002/NHSPS. 1.9 Safe disposal of waste (including sharps) Health Technical Memorandum (HTM 07-01) contains the regulatory waste management guidance for all health and care settings (NHS and non-NHS) in England and Wales including waste classification, segregation, storage, packaging, transport, treatment and disposal. Health and Safety (Sharp Instruments in Healthcare) Regulations 2013 outline the regulatory requirements for employers and contractors in the healthcare sector in relation to the safe disposal of sharps. Definitions Healthcare (including clinical) waste: Clinical waste means waste from a healthcare activity (including veterinary healthcare) that: contains viable micro-organisms or their toxins which are known or reliably believed to cause disease in humans or other living organisms. For example, if a patient is known or suspected to be infected, or colonised by an infectious agent. Clinical judgement should be applied in the assessment of waste and should consider the infection status of a patient and the item of waste produced. contains or is contaminated with a medicine that contains a biologically active pharmaceutical agent, or is a sharp, or a body fluid or other biological material (including human and animal tissue) containing or contaminated with a dangerous substance within the meaning of Regulation (EC) No 1272/2008 of the European Parliament and of the Council on classification, labelling and packaging of substances and mixtures, as amended from time to time. Offensive waste is waste that: is not clinical waste, is not infectious, but may contain body fluids, secretions or excretions, is non-hazardous, and falls within waste codes 18 01 04 if from healthcare, or 20 01 99 if from municipal sources. Table 1: Categories of waste and segregation at source Category Segregation Treatment/Disposal Offensive (non-infectious) Yellow bag with black stripe (tiger) bag Energy from waste, landfill or other permitted processes Clinical waste (infectious only) UN approved orange bag, UN approved box or sharps container For alternative treatment Healthcare waste contaminated with non-hazardous pharmaceuticals or chemicals) UN approved yellow bag, UN approved box or sharps container For incineration or other permitted process Waste contaminated with cytotoxic or cytostatic medication UN approved purple bag, UN approved box or sharps container For incineration Non-hazardous pharmaceuticals (no sharps) Blue box/container For incineration or other permitted process Anatomical waste/full blood bag and blood preserves UN approved red lidded container For incineration only Domestic Black/clear bags Energy from waste, recovery or landfill Recycling Clean, green or other colour bag Recycling Safe waste disposal at care area level: Always dispose of waste: immediately and as close to the point of use as possible; and into the correct segregated colour coded rigid container or sharps box if a sharp Liquid waste, eg, suction canisters, must be rendered safe by adding a polymer gel or compound to the container prior to placing in the orange lidded leak proof bin or yellow lidded leak proof bin if contaminated by pharmaceutical waste bags must be no more than 2/3 full and no more than the UN approved weight and must be securely tied using a plastic tie or secure knot using a 'swan neck' to close. Waste must be traceable back to ward/care area or department, this may be achieved by writing on bags (prior to use), attaching sticky labels or uniquely numbered tags with the post code on them. store all waste in a designated, safe, lockable area while awaiting collection. Collection schedules must be acceptable to the care area and there should be no build-up of waste receptacles. Local guidance on management of waste at care level, eg, domiciliary settings should be followed. Sharps containers (for safety devices, refer to section 1.10) Sharps containers must: have a handle (small community boxes do not require a handle) and temporary closure mechanism, employed when box is not in use be disposed of when the manufacturers' fill line is reached be labelled with point of origin and date of assembly and disposal. Where re-usable sharps containers are used, organisations must have a protocol in place to assure themselves of safe use and reprocessing. Further information can be found in the Health Technical Memorandum (HTM 07-01). 1.10 Occupational safety: prevention of exposure (including sharps injuries) The Health and Safety (Sharp Instruments in Healthcare) Regulations 2013 outline the regulatory requirements for employers and contractors in the healthcare sector in relation to: arrangements for the safe use and disposal of sharps; provision of information and training to employees; investigations and actions required in response to work related sharps injuries. There is a potential risk of transmission of a BBV (blood borne virus) from a significant occupational exposure and staff must understand the actions they should take when a significant occupational exposure incident takes place. There is a legal requirement to report all sharps injuries and near misses to line managers/employers. A significant occupational exposure is: a percutaneous injury eg injuries from needles, instruments, bone fragments, or bites which break the skin (abrasions, cuts, eczema, etc); and/or exposure of mucous membranes including the eye From splashing of blood or other high risk body fluids. For the management of an occupational exposure incident see Appendix 10. Safety devices Health and Safety (Sharp Instruments in Healthcare) Regulations 2013 are concerned with reducing and eliminating the number of 'sharps' related injuries which occur within healthcare. Its basic guidance is: avoid unnecessary use of sharps if use of medical sharps cannot be avoided, source and use a 'safer sharp' device; if a safer sharp device is not available then safe procedures for working with and disposal must be in place eg sticky mats, sharps bins, safety procedures and training. Sharps handling must be assessed, kept to a minimum and eliminated, if possible, with the use of approved safety devices, manufacturers' instructions for safe use and disposal must be followed needles must not be re-sheathed/recapped or disassembled after use sharps must not be passed directly hand to hand used sharps must be discarded at the point of use by the person generating the waste always dispose of needles and syringes as 1 unit if a safety device is being used safety mechanisms must be deployed before disposal. When transporting sharps boxes for community use these must be transported safely with the use of temporary closures. Further information can be found in occupational exposure management (incl. sharps) literature review. > The national infection prevention and control manual (NIPCM) is an evidence-based practice guide for use in England. It contains standard infection control precautions (SICPs) and transmission-based precautions (TBPs), which when applied correctly can help reduce the risk of healthcare associated infection (HCAI) and ensure the safety of those being cared for, staff and visitors in the care environment. The NIPCM aims to: make it easy for health and care staff to apply effective infection prevention and control (IPC) precautions reduce variation and optimise infection prevention and control practices throughout England help reduce the risk of HCAI help align practice, monitoring, quality improvement and scrutiny. The NIPCM should be applied by all NHS staff involved in patient care. Furthermore, the principles in the NIPCM should be applied across all care settings (including acute, community and social care), complementing specific guidance produced for these settings, complementing specific guidance produced for these settings. Several supporting tools (appendices) are available to complement the NIPCM. The NIPCM is underpinned by scientific literature reviews, these are primarily aimed at infection control specialists and summarise the available evidence, highlight any research gaps. Currently, the NIPCM for England uses the Scottish NIPCM literature reviews as its evidence base. NHS England will use these reviews for an interim period, during which we will develop the IPC evidence base for England. This guidance sets out the defined methodology that will be used to achieve this aim. It is fundamental to the integrity and applicability of the NIPCM that a wide group of stakeholders is involved in its development. Involving experts from all appropriate multidisciplinary groups during development of the NIPCM, its associated literature reviews and supporting tools ensures its recommendations are evidence-based, appropriate, practical and acceptable in all healthcare settings. Two standing groups participate in developing the NIPCM: the NIPCM consensus group and the NIPCM clinical oversight group. In addition, topic-specific 'review' groups will be formed to support NIPCM literature reviews. Together, the NIPCM review, consensus and clinical oversight groups form the NIPCM guideline development group. The membership of the NIPCM consensus group is intended to reflect IPC practitioners at the front line across a range of services (acute, community, mental health, etc). All members must be employed in a relevant position, ie related to HCAI and infection control. The NIPCM consensus group membership is formed of frontline IPC specialists with responsibility for implementing IPC policy and guidance. The group has representation from across the seven geographical regions including acute, community, mental health and learning disability, and ambulance specialties. The full membership is available in the consensus group terms of reference. The NIPCM clinical oversight group has a wider professional membership. The following organisations and specialties are represented (or included for engagement/consultation purposes) on the NIPCM consensus group: Healthcare Infection Society (HIS) Infection Prevention Society (IPS) Royal College of Nursing Academy of Medical Royal Colleges Occupational medicine NHS England regional IPC leads UK Health Security Agency (UKHSA) Health and Safety Executive (HSE) Chartered Physiotherapy Society Care Quality Commission (CQC) NHS England Primary care services (including dentistry) Department of Health and Social Care (DHSC). NB This list is not exhaustive as membership is reviewed quarterly refer to Terms of Reference for current membership. Lay representatives are also engaged for the NIPCM clinical oversight group. They will have an interest in the NHS and in reducing the incidence and impact of HCAI through applicable and accessible infection prevention and control guidance. The lay representatives are expected to attend all meetings and comment on recommendations from the perspective of a patient or visitor. A specialist clinical 'review' group will also be formed for NIPCM literature reviews. The membership of these groups will vary but will comprise topic-specific (clinical and academic) experts and individuals with expertise in literature reviewing and guideline development. Members may also be drawn from the consensus and clinical oversight groups with relevant expertise. A key role of the groups involved in the NIPCM development is to represent the views of all appropriate staff members/groups within and across the represented organisation/regions. The roles and responsibilities of all members of the review, consensus and clinical oversight groups are laid out in full in the terms of reference corresponding to each group. Briefly, members must: contribute to the consultation process on the NIPCM (including literature reviews and any supporting documents/tools), feeding back the views of the professional groups/organisations they represent, such as barriers to implementation contribute to identifying evidence/research gaps in the literature pertaining to the NIPCM and support the development of research studies to enhance the evidence base. Review group members will provide specialist input into the review process. NHS England scientists will lead on developing research questions, search strategies, identification of evidence, critical appraisal/data extraction, summarising evidence and final write-up. Review group members will provide expert input and critique/peer review at each of these stages. Review group members will provide additional clinical or scientific advice and input to the review by assessing: the technical or clinical validity and clarity of research and identifying any omitted terms, acronyms or synonyms the appropriateness of the eligibility criteria developed the studies identified by NHS England for full text review to identify any additional studies not identified or erroneously excluded the final draft of the literature review as part of the defined stakeholder engagement process (peer review). Review group members are not directly involved in the eligibility assessment, data extraction, appraisal, or in the drafting of the review. Meetings of both the consensus and clinical oversight groups are scheduled on a quarterly basis. For a meeting to be quorate at least 50% of members (including the chair or their deputy) must be present. All members (including chairs) of the NIPCM review, consensus, and clinical oversight group are required to declare any competing interests (see Appendix 2). NHS England literature reviews that inform the NIPCM are produced using systematic (iterative) methods that are conducted in accordance with current best practice, eg Cochrane review methodologies. The choice of literature review design is based on the topic, available resources and timescales required and must be agreed by the clinical oversight group and any review group formed for the purpose of conducting the review. Overview and intended use: Used when a systematic review is likely to be too restrictive to meet research objectives; for example, to clarify concepts or definitions, assess or map a body of evidence or identify knowledge gaps. Not typically intended for the development of evidence-based recommendations but this may be achievable with thoughtful design. Rigour of methodology is similar to a systematic review (ie an iterative and structured approach). Conducted in accordance with Joanna Briggs Institute methodology and reported in accordance with PRISMA extension for scoping reviews. No. of reviewers (NHS England scientists): 2 Evidence sources: Purposefully broad, includes grey literature, policy and legislation, other guidance/ guidelines, and multiple biomedical databases are searched. May include hand searching and snowball searching. Screening of search results: Screening of literature search results is performed in full by two reviewers. Critical appraisal and grading of evidence: Typically, no formal critical appraisal or grading of evidence is performed; data is extracted into evidence tables and checked (at least 30%) by the second reviewer. Evidence synthesis and reporting: A narrative synthesis is performed in accordance with SwIM and results are reported using the PRISMA extension for scoping reviews (PRISMA-SCR). Overview and intended use: Used to answer a research question that is extremely focused or to examine a topic where little is known/had been published or to examine a highly specialised and focused topic/specialty. Uses an explicit, systematic (iterative) and structured process to develop evidence-based recommendations. Similar to a systematic review but with a more restricted scope and typically shorter timeframe. Literature searches are restricted in number, and eligibility criteria are tighter/more focused than in a systematic review (eg, in terms of eligible date range). Conducted and reported in accordance with best practice (Cochrane and PRISMA). No. of reviewers (NHS England scientists): 2 Evidence sources: Purposefully focused; omits grey literature unless identified as key evidence source by the review group; typically a single biomedical database is searched Screening of search results: Screening of literature search results is performed in full by a single reviewer; a second reviewer checks a minimum 20% sample during the first screen and 100% of excluded studies from the second screen. Critical appraisal and grading of evidence: Varies by timescale and topic and is agreed by the review group in advance. Evidence synthesis and reporting: Typically, a single evidence table without an evidence summary reported in accordance with PRISMA and SwIM. Timescale: At least 3 months Overview and intended use: Used to answer focused research question(s) on a specific topic to develop evidence-based recommendations. Uses an explicit, systematic (iterative) and structured process to identify, select and critically appraise evidence, typically from primary research. Conducted and reported in accordance with best practice (Cochrane, PRISMA, AGREE). No. of reviewers (NHS England scientists): 2 Evidence sources: Multiple biomedical databases are searched. Includes grey literature, policy and legislation, other guidance/guidelines. Screening of search results: Two reviewers independently screen all search results against the eligibility criteria. Critical appraisal and grading of evidence: Critical appraisal and data extraction of all evidence as detailed in section 7 is performed by a single reviewer. A second reviewer performs a minimum 30% check. Evidence synthesis and reporting: Evidence tables and evidence summaries produced for each research question by a single reviewer. These are checked by the second reviewer and peer reviewed by the review group. Reported in accordance with PRISMA and SwIM. Overview and intended use: Used to summarise and collate mandatory recommendations or directives from existing legislation or policy, nationally and internationally recognised evidence-based guidelines. This may be used in combination with the two person systematic review if some of the research objectives require searching biomedical databases and appraising peer-reviewed published scientific literature etc. Follows a similar process to a two-person systematic review but with a single reviewer identifying eligible evidence and performing data extraction. No. of reviewers (NHS England scientists): 1 Evidence sources: Focused on current national policy and legislation, and up to date (