

| 可能造成風險暴露的體液 | 可能造成風險暴露的體液 |
|---|---|
| 可能具有感染性 CSF Pleural fluid Pericardial fluid Peritoneal fluid Amniotic fluid Synovial fluid Tissue samples | 不具有感染性 (如果沒看到血) Tears Saliva (except in dental setting) Urine Feces Sweat Emesis |
| | 風險暴露如何發生 |
| | Exposures occur through |
| 工作上的風險暴露 | Needlesticks or cuts from other sharp instruments contaminated with an infected patient's blood Or Contact of the eye, nose, mouth, or skin with a patient's blood. |
| | |
| 風險暴露有哪幾種狀況? | 風險暴露有多嚴重? |
| Exposures known to pose a risk of transmission for bloodborne pathogens: – Percutaneous injury (hollow needle> solid sharp) – Splash on mucous membrane – Splash on non-intact skin The risk of transmission increases with larger volumes of fluid and more severe injuries | Difficult to asses: up to 70% of exposures go unreported 1990 estimate: 500,000 exposures/year Impossible to measure the psychological stress that an exposure places on a health care worker |
| | Marcus, R. et al. Ann Emerg Med 1995;25:77 Henry, K. Minnesota Medicine 1995;78:41-44 |

| 風險 | 暴露 | 的愿 | 太 染オ | 機會 |
|----|----|----|-------------|----|
| | | | | |

| Hepatitis | Outbreak or Needlestick Exposure Transmission Rate (%) | Prevention | Comment |
|-----------|---|---|---|
| A | 10-30 | Vaccine not given routinely to HCWs Immune globulin in outbreak setting | ACIP advises vaccine "is or might be" indicated |
| B | | HBV vaccination | HBV vaccination acceptance for dialysis patients (58%) and staff (58%) has increased in recent years |
| cAg | 3 | HBIG if appropriate | |
| eAg* | 20-40 | HBV vaccination HBIG if appropriate | |
| с | 1-10 | Immunoglobulin not recommended Interferon therapy for acute disease appears promising | Prevalence in U.S. dialysis units: 8.4% (patients) and 1.7% (staff) |
| Delta | Unknown rate; outbreaks de- scribed only in dialysis units | HBV vaccination | Segregate HBsAg-positive dialysis patients by delta antibody status |
| E | None described | Unknown | Probably no increased seroprevalence among dialysis or other patients |

ACIP, Advisory Committee on Immunization Practices: Ag. antigen: HBV, hepatitis B virus; HBIG, hepatitis B immune globalin; HBsAg, hepatitis B surface antigen.

醫療照顧者不可不知

- Most exposures do not result in infection.
- Following a specific exposure, the risk of infection may vary with factors, including:
 - The pathogen involved
 - The type of exposure
 - The amount of blood involved in the exposure
 - The amount of virus in the patient's blood at the time of exposure



陽轉率(Pre-HAART)

| | All Patients | | HIV-1 infected Patients | | |
|-----------------|--------------|-----------------|-------------------------|-----------------|---|
| | HCW report | Total Exposures | HCW report | Total Exposures | |
| Blood | 309 | 10,008 | 136 | 2,712 | 0 |
| Sputum | 112 | 3,144 | 47 | 804 | 0 |
| Urine | 155 | 3,780 | 61 | 912 | 0 |
| Feces | 49 | 828 | 20 | 300 | 0 |
| Other fluids | 93 | 3,096 | 40 | 840 | 0 |
| Total | 337 | 20,856 | 149 | 5,568 | 0 |

職業暴露的風險

- Of 5 868 samples, 1.1 % had reactive or borderline ELISA test
- Of 1 344 participants, 2.6% had positive ELISA, 2.1% false positive, and 7 * had positive.
- Transmission 1 case per 179 exposures. (0.56% chance per exposure, Cl:0.01%~3.06%)
- Different exposure risk
 Mucous membrane 0.86%, Cutaneous exposure 0.11%
- Overall occupational risk 0.3%

Annals of Internal Medicine 1990;113:740-46

處理風險暴露



時間就是病毒

- Animal studies show that PEP should be given within 2-8 hours of exposure for maximal effect
- PEP may have some benefit up to 36 hrs but seems to be ineffective if given later

JID 1991;163:625 JID 1993;168:825

HIV:風險暴露後的追蹤

- 規則的追蹤時間 (eg 6,12 weeks and 6 months)
- Testing should continue for 12 months if the HCW contracts HCV from the exposure
- Unclear if testing should be prolonged in exposures to pts with HIV and HCV or in HCW who have history of impaired Ab responses

HIV:風險暴露後的諮詢

- For 3 months following exposure HCW should avoid
 - Unprotected sex
 - Pregnant
 - Donate blood, organs, or semen
 - Sharing razors, toothbrushes
- HCW should consider stopping breast feeding (risk of perinatal transmission and drugs may get into breast milk)

HIV暴露後次級預防

| | | 暴露来源病患 | 的 HIV 感染狀態 | | |
|------------------------|---------------------------------|--------------------------|---|--|----------|
| 暴露的種類 | 第一規 (Class 1) 感染状態 | 第二級 (Class 2) 成染狀態 | 病患的感染狀態 不詳 | 不知來源病患 | 未感染 HIV |
| 較不嚴重者 ⁽¹⁸⁾ | 建議使用 基本 PEP ^(国王) | 建議使用 加強 PEP | 通常並不須要使 用 PEP:但若來 源病患有感染 HIV 的危險性 時,可考慮使用 基本 PEP | 通常並不須要使 用 PEP:但若推 測可能的來源病 患有感染 HIV 的 危險性時,可考 慮使用基本 PEP | 不需使用 PEP |
| 比較嚴重者 ^(11二) | 建築使用 加強 PEP ^(IIN) | 建镁使用 加強 PEP | 通常並不須要使 用 PEP:但若來 源病患有感染 HIV 的危險性 時,可考慮使用 基本 PEP | 通常並不須要使 用 PEP: 但若推 測可能的來源病 患有感染 HIV 的 危险性時,可考 慮使用基本 PEP | 不需使用 PEP |



次級預防用藥

□ abacavir (ABC) + lamivudine $(3TC^{B})$ or as fixed dose combination

五、 efavirenz (Stocrit[®]; EFV)(在已知懷孕或在生育年齡的婦女要注意致

一、 lopinavir/ritonavir (Kaletra[®]複方; LPV/RTV)(優先選用之配方)

-、 zidovudine + lamivudine (Combivir[®]複方)(優先選用之配方)

(Kivexa[®]複方)(需特別注意過敏的可能^(nx))。

Ξ · indinavir (Crixivan[®]; IDV) + ritonavir (Norvir[®]; RTV)

= · lamivudine + didanosine (Videx[®] EC) = · lamivudine (3TC[®]) + stavudine (d4T)

∴ atazanavir (Reyataz®; ATV)

四、 nelfinavir (Viracept[®]; NFV)

畸胎的可能)

