Audit of the use of drotrecogin alfa (activated) for severe sepsis (Xigris)

| TO | Emma North  
| Fax Number: | 020 7388 3759 |

**ADMISSION**

| CMP Admission Number: |  
| Date of admission to your unit: |  

**INFUSION**

| Date of first infusion of Xigris: |  
| Time of first infusion of Xigris: |  

Did the patient receive the complete 96-hour infusion of Xigris?  
If **No**, please state why not:

Was there any interruption in the infusion of Xigris?  
If **Yes**, on what date and at what time did the interruption occur?  
What was the duration of the interruption?  
What was the reason for the interruption in the infusion?  

Was this infusion given as part of a study/trial?  
If **Yes**, please indicate the site:

**INFECTION**

Do you know the primary site of this patient's infection?  
If **Yes**, please indicate the site:  
Lung:  
Abdomen:  
Urinary tract:  
Other:  
If **Other**, please state site:

Was a blood culture done prior to the infusion of Xigris?  
If **Yes**, was it positive?  
Was an organism cultured for this patient's infection?  
If **Yes**, please tick all organisms cultured:

Gram-negative bacteria  
Gram-positive bacteria  
Fungus  
Name of primary pathogen (or state if not known)  

**ADVERSE EVENTS**

Did a serious adverse event occur (any adverse event assessed as serious by the clinician)?  
If **Yes**, was it a: Serious bleeding event?  
Thrombotic event?  
Other event?  
If a **serious bleeding event**, please indicate site(s):  
Gastrointestinal bleed:  
Genitourinary bleed:  
Skin or soft tissue bleed:  
Intrathoracic bleed:  
Intraabdominal bleed:  
Other (source unidentified) bleed:  
Intracranial bleed:  
Retroperitoneal bleed:  
Other (source identified) bleed:  
If **Other (source identified) bleed**, please state site(s):

---

§ Serious bleeding events are defined as any intracranial haemorrhage, any life-threatening bleed or any bleeding event requiring the administration of >= 3 units of packed red blood cells per day for two consecutive days, or any bleeding event assessed as serious by the clinician.