Sepsis scoring systems and use of the Sepsis six care bundle in maternity hospitals

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Abstract

Background: This study aimed to assess the predictive power of three different Sepsis Scoring Systems (SSSs), namely maternity Systematic Inflammatory Response Syndrome (mSIRS), quick Sepsis-related Organ Failure Assessment (qSOFA) and Modified Early Warning System (MEWS) in identifying sepsis by comparing them with positive culture. This study also sought to evaluate compliance with using the Sepsis Six Care Bundle (SSCB) operated in an individual health board.

Methods: A retrospective cohort study was conducted in 3 maternity hospitals of a single Scottish health board that admitted 2690 pregnancies in a 12 weeks period in 2016. Data for study was obtained from medical notes, handheld and electronic health records for women who were prescribed antibiotics with a confirmed or suspected diagnosis of sepsis. Data on clinical parameters was used to classify women according to mSIRS, qSOFA and MEWS as having sepsis or not and this was compared to results of positive culture to obtain sensitivity, specificity, positive predictive value (PPV), negative predictive value (NPV) and area under Receiver Operating Characteristic curve (AUROC) along with their 95% confidence intervals. Data was also obtained on SSCB compliance.

Results: A total of 89 women were diagnosed with sepsis, of which 14 had missing data, leaving 75 for final analysis. Sensitivity, specificity, PPV, NPV and AUROC of mSIRS and MEWS were almost similar with AUROC of both being around 50%. Only 33 (37.1%) had identifiable sepsis six sticker displayed on medical notes and only 2 (2.2%) had all elements of SSCB delivered within the recommended one-hour post-diagnosis period. Blood culture and full blood count with other lab tests had been performed for most women (97%) followed by intravenous antibiotics and fluids (93.9%).

Conclusions: mSIRS and MEWS were quite similar in detecting sepsis when compared to positive culture, with their ability to detect sepsis being close to chance. This underlines the need for creating a valid SSS with high sensitivity and specificity for clinical use in obstetric settings. Clinical use of SSCB was limited despite it being a health board policy, although there is considerable possibility of improvement following detailed audits and removal of barriers for implementing SSCB.

Keywords: Sepsis, Sepsis scoring systems, Sepsis diagnosis, Obstetric, Maternity, Systemic inflammatory response syndrome, SIRS, qSOFA, MEWS, Early warning systems, Sepsis six care bundle

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Background

Sepsis was initially defined at the 1991 ACCP/SCCM Consensus Conference as “a host’s systemic inflammatory response syndrome (SIRS)” [1], based on a score obtained using the patient’s temperature, heart rate (HR), white cell count (WCC) and respiratory rate (RR) [1, 2]. In an obstetric population, changes associated with pregnancy and labour can make the SIRS baseline parameters misleading for the identification of sepsis [3]. These alterations in a woman’s normal physiology can mask the initial phase of sepsis and delay diagnosis, hindering appropriate clinical intervention and subsequent recovery [4]. Interpretation of other biomarkers of infection such as C-reactive protein (CRP) and WCC can also vary with the mode of delivery that women experience [5, 6]. As a consequence, a number of alternate Sepsis Scoring Systems (SSSs) were developed for maternity use such Sepsis in Obstetric Score (SOS) [3] or others such as maternity SIRS (mSIRS) [7] which are in use only in certain medical systems.

Not only for obstetric cases but even for the normal population, a re-evaluation of the definition of sepsis and SIRS after 24 years of clinical use found it to be deficient in both specificity and sensitivity, as it failed to capture many true cases of sepsis and identified infections that are not necessarily sepsis [1]. The re-evaluation indicated that the SIRS construct, although useful in identifying patients with infection, had limited specific applicability to sepsis [1]. The definition of sepsis was overhauled during the 3rd International Consensus Definition for Sepsis and Septic Shock [1]; the new definition of sepsis being “life-threatening organ dysfunction caused by a dysregulated host response to an infection” [1]. In order to detect sepsis in line with the new definition, a Sepsis-related Organ Failure Assessment (SOFA), and a quick SOFA (or qSOFA) were created, comprising only three indices: mental status, systolic blood pressure (SBP) and RR [1]; although SOFA or qSOFA may not capable of replacing SIRS [2].

Apart from variations of SIRS, SOFA and qSOFA, the Royal College of Obstetricians and Gynaecologists UK, endorsed a pregnancy-specific scoring system, the Modified Obstetric Early Warning Score (MOEWS) [3, 8] which was related to Modified Early Warning Score (MEWS) for non-obstetric population [9]. A number of MOEWS are in use in various facilities globally but very few have been validated [9]. Evidence suggests that most of these MOEWS are not as good when compared to MEWS for obstetric use [9].

In addition to developing various SSSs for early detection of sepsis during obstetric care, it was also essential to recognize that patients’ deterioration can be limited by the early management of sepsis. This has led to the application of care bundles as a mechanism for minimizing harm and enhancing patient care [10]. The Sepsis Six Care Bundle (SSCB) was recently introduced into the maternity hospitals of the health region under examination in this study, in the form of a sticker on patients’ notes, with the aim of delivering all six elements of the bundle (Additional file 1: Sepsis Six Sticker) within 1 h of sepsis being provisionally diagnosed. However, compliance has not been uniform and in line with the suggestions of SSCB [11].

Thus, there are many variations of SSSs in obstetric use across the world and very few have been validated including some we were interested in. Hence, we assessed mSIRS (since this was recommended by the health board where this study was conducted), qSOFA (an improvement over the earlier SIRS) and MEWS (which performed better than all MOEWS [9]) in identifying sepsis by applying these criteria to all pregnant women treated for sepsis and by comparing them to positive culture as a gold standard. In addition, we also evaluated compliance with using the SSCB.

Methods

Study design and settings

A retrospective observational cohort study was conducted within three maternity hospitals situated in a single Scottish health region that admitted 2690 pregnant women in a 12 weeks period in 2016. The study sample comprised all women admitted to these maternity hospitals during the study period who received antibiotic therapy for a suspected or confirmed diagnosis of sepsis.

Data collection and subject identification

Patients were identified through hospital handover sheets or/and drug Kardexes® (records of patients’ medication prescription and administration). Women ≤16 years of age were excluded. Women were also excluded if antibiotics were prescribed prophylactically; such women were included only if they required additional treatment following a diagnosis of sepsis.

Individual demographic characteristics including patient’s age, body mass index (BMI), gestational age, mode of delivery and unit of hospitalization were collected. Data on each patient’s temperature, heart rate (HR), respiratory rate (RR), systolic blood pressure (SBP) and mental status were collected from individual MOEWS charts, while individual data on WCC and microbiology reports (positive culture) were collected from patient electronic health records (EHRs).

Patient’s clinical notes were then reviewed to determine whether the SSCB had been commenced and used appropriately. This evaluation was not reported for patients who were commenced on the bundle without the use of the SSCB sticker.
### Table 1 Demographic information of pregnant women (N = 89) diagnosed with suspected sepsis in maternity wards of 3 hospitals of a health board in Scotland

| Demographic variables | Values (N = 89) |
|-----------------------|----------------|
| **Age**<sup>a</sup> | Mean ± SD (Min-Max) 29.8 ± 5.3 (19-45) |
| **BMI** | Normal (18.5–24.9) 36 (46.7%) |
| Overweight (25–30) | 22 (28.6%) |
| Obese (≥ 30) | 19 (24.7%) |
| **Gestational age**<sup>b</sup> | First trimester 0 |
| Second trimester | 5 (6%) |
| Third trimester | 79 (94%) |
| **Delivery mode**<sup>c</sup> | Emergency caesarean section 41 (46.1%) |
| Spontaneous vaginal delivery | 22 (24.7%) |
| Instrumental delivery | 14 (15.7%) |
| Elective caesarean section | 1 (1.1%) |
| **Unit of hospital admission** | Antenatal or Postnatal ward 80 (89.9%) |
| High Dependency Unit | 7 (7.9%) |
| Intensive Care Unit | 2 (2.2%) |

<sup>a</sup>Data were missing for 12 women; <sup>b</sup>Data were missing for 5 women; <sup>c</sup>Data were missing for 11 women, of whom 7 were diagnosed with sepsis in the antenatal period

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### Data management

Based on the various SSSs that were evaluated, the values of all required patient parameters such as HR, RR, SBP, mental status etc. were dichotomized to be normal or abnormal. These dichotomized values were then combined to create a composite score for a particular SSS to evaluate whether a patient had sepsis or not based on a particular SSS. Positive culture results were used to evaluate the selected SSSs.

### Data analysis

All analysis was performed using SPSS Statistics for Windows, Version 23.0 (Armonk, NY: IBM Corp.). Means, standard deviations and range was calculated for all continuous variables. Counts and percentages were calculated for all categorical variables. We evaluated three SSSs – mSIRS, qSOFA and MEWS (we selected MEWS and not any of the MOEWS since a recent study discovered MOEWS to be less effective than MEWS in detecting sepsis early in obstetric settings [9]) with positive culture results as gold standard to obtain sensitivity, specificity, positive predictive value (PPV) and negative predictive value (NPV) along with their 95% confidence intervals (CIs). To summarize our evaluation of SSSs we calculated Receiver Operating Characteristic (ROC) curve along with area under the ROC (AUROC) and 95% CIs. We also assessed compliance with the SSCB in our study by calculating the counts and percentages of patients receiving the six elements of SSCB.

### Results

This study included a total of 89 (3.3%) women, diagnosed with sepsis, from a total of 2690 pregnancies, in a single health region in Scotland during a 12-week study period in 2016.

Table 1 summarises the demographic details of the 89 women based on the data available in the EHRs. The average age of these women was 29.8 (± 5.3) years and ranged from 19 to 45 years. A total of 19 (24.7%) women were classified as obese with Body Mass Index (BMI) equal to or greater than 30, with most women (46.7%) having a normal BMI. Most women (94%) were in the third trimester, with the greatest proportion of deliveries occurring via emergency caesarean section (46.1%), followed by spontaneous vaginal delivery (24.7%). The majority of women (89.9%) were admitted either into the antenatal or postnatal wards with very few in high dependency unit (7.9%) or intensive care unit (2.2%).

Data for evaluating Sepsis Scoring Systems (SSSs) was available only for 75 out of 89 (84.3%) women. Various Sepsis Scoring Systems, namely mSIRS, qSOFA and MEWS (Table 2), were compared with positive culture (Table 3), to obtain characteristics such as sensitivity, specificity, PPV, NPV and AUROC. mSIRS and MEWS were found to be comparable with regards to most characteristics – sensitivity [mSIRS: 0.655 (95% CI – 0.643-0.668); MEWS: 0.690 (95% CI – 0.678-0.702)], specificity [mSIRS: 0.391 (95% CI – 0.379-0.404); MEWS: 0.304 (95% CI – 0.292-0.316)], PPV [mSIRS: 0.404 (95% CI – 0.391-0.417); MEWS: 0.385 (95% CI – 0.372-0.397)], and NPV [mSIRS: 0.643 (95% CI – 0.630-0.655); MEWS: 0.609 (95% CI – 0.596-0.621)]. AUROC for both were close to 50% (mSIRS AUROC – 0.524 (95% CI - 0.388-0.659) and MEWS AUROC – 0.497 (95% CI - 0.354-0.639)). Thus, specificity, PPV, NPV and AUROC were slightly higher for mSIRS compared to MEWS. qSOFA on the other hand was positive (score ≥ 2) for only 2 women both of whom had negative cultures (Table 3).

We also evaluated the use of Sepsis Six Care Bundle (SSCB) for 33 (37.1%) women (Table 4), who had the sepsis six sticker on their medical notes, and discovered that most women received 5 of the 6 elements of SSCB. Most women had blood culture, and full blood count with other lab tests performed (97%) followed by intravenous antibiotics and intravenous fluids (93.9%). Catheterization was done for 78.8% women and 27.3% women administered oxygen. It is important to note here that only 2 cases received all 6 elements of SSCB.
within the prescribed 1 h of being diagnosed with sepsis – at 35 and 40 min from time zero.

**Discussion**

Our study on 89 women, in a single health board in Scotland, which compared three different SSSs – namely mSIRS, qSOFA and MEWS, with positive culture, revealed comparable performance by mSIRS and MEWS with similar values for sensitivity, specificity, PPV, NPV and AUROC. Most importantly the values for AUROC for both mSIRS and MEWS were around 50% indicating that their ability to predict sepsis was close to chance compared to positive culture. Estimates for qSOFA were unreliable in our study due to low number of women with positive qSOFA.

On evaluating compliance with the use of SSCB, only 33 (37.1%) of these women were found to have sepsis six sticker on their medical notes denoting low levels of compliance with providing timely recommended sepsis care. Among those who received SSCB, only 2 women received it within the recommended 1 h following diagnosis. While most received 4/6 elements of SSCB, only around 79% received catheterization and the minority (27.3%) were administered oxygen.

Our findings regarding comparison of SSSs with positive culture threw up meaningful results only for mSIRS and MEWS. mSIRS is a modification of the SIRS criteria and has been in use in the hospitals of the health board since 2015 [7]. mSIRS performance was similar to that of MEWS which indicated that mSIRS was probably one of the better performing SSS in obstetric settings. This can be inferred in the light of findings where MEWS was compared to six different published MOEWS and was found to perform better than all of them [9]. The different MOEWS were reported to have low PPV, ranging between 1.4 and 5.1%, while AUROC ranged from 0.52 to 0.72 [9].

The use of SIRS criteria in the original definition of sepsis were not specific and its values can be elevated in cases of other non-infectious disease such as burns or injury and in cases unrelated to infection [12, 13]. In addition, women undergo changes in their physiological and laboratory measurements during pregnancy and post-partum periods. These changes overlapped with the SIRS criteria and made the diagnosis critical, particularly when respiratory rate, heart rate, white cell count and blood pressure values, expected to be outside the adult normal reference range [12, 13].

An important thing to note here is the continued use of mSIRS (though SIRS was found highly inadequate after decades of use [1]) in the health board that was studied and the presence of numerous MOEWS in various obstetric setting across the world; this points towards the lack of consensus [13] about a single validated SSS which is needed for use in obstetric settings and is currently missing.

**Table 3** Comparison of Sepsis Scoring Systems against Positive Culture \(^a\) for pregnant women (\(N = 75\)) diagnosed with probable sepsis in maternity wards of 3 hospitals of a health board in Scotland

| Characteristics | Sepsis Scoring Systems for Maternity Care % (95%CI) \(^b\) | Maternity SIRS | qSOFA \(^c\) | MEWS |
|-----------------|-------------------------------------------------|----------------|-------------|-------|
| Sensitivity     | 0.655 (0.643–0.668)                              | 0.690 (0.678–0.702) |              |       |
| Specificity     | 0.391 (0.379–0.404)                              | 0.957 (0.951–0.962) | 0.304 (0.292–0.316) |       |
| PPV             | 0.404 (0.391–0.417)                              | 0.385 (0.372–0.397) |              |       |
| NPV             | 0.643 (0.630–0.655)                              | 0.603 (0.589–0.603) | 0.609 (0.596–0.621) |       |
| AUROC           | 0.524 (0.388–0.659)                              | 0.301 (0.045–0.558) | 0.497 (0.354–0.639) |       |

\(^a\)29 patients had a positive culture; \(^b\) CI: Confidence Interval; \(^c\) Only 2 patients had qSOFA score of 2 and both had negative cultures
partum period during pregnancy, childbirth, post-abortion, or the post-
defined as organ dysfunction resulting from infection
ous biomarkers due to the non-specific or non-sensitive
early identification tool is necessary so that treatment
after therapy had been initiated for these women, and an
culture results [16]. These shortcomings with cul-
which are associated with a delay in administering the
appropriate therapy and management to these women [18]. Culture-proven sepsis results were obtained 48 h
after therapy had been initiated for these women, and an
early identification tool is necessary so that treatment
can be delivered early. The limited applicability of vari-
ous biomarkers due to the non-specific or non-sensitive
nature of these criteria, arising from the altered

| Care/therapy                        | Compliance with the care at any time point n (%) |
|-------------------------------------|-----------------------------------------------|
| Oxygen                              | 9 (27.3%)                                     |
| Blood culture                       | 32 (97%)                                      |
| Full blood count and other lab tests| 32 (97%)                                      |
| Intravenous antibiotics             | 31 (93.9%)                                    |
| Intravenous fluid                   | 31 (93.9%)                                    |
| Catheter                            | 26 (78.8%)                                    |

Note: Only for 2 cases SSCB was delivered within the prescribed one-hour period - reported times being 35 and 40 min from time zero

It has been proposed that the qSOFA definition should be able to identify women at an early stage of severe ma-
three SSSs show uncertainty about systems to be used in diagnosis and
culture-proven sepsis is still the diagnostic gold standard. There were only 29 positive culture results among
women (38.6%), which is in line with the evidence [16]. Despite being the gold standard there are severe
deficiencies in using culture tests in clinical settings due to poor specificity and precious time lost in waiting for
culture results [16]. These shortcomings with cul-
tests further underline the need for creating an ef-
effective SSS with high sensitivity and specificity for early
use in clinical settings.

The early identification and management of sepsis in
maternity is recommended [17], as delay in diagnosis or
treatment can lead to maternal mortality and morbidity, which are associated with a delay in administering the
appropriate therapy and management to these women [18]. Culture-proven sepsis results were obtained 48 h
after therapy had been initiated for these women, and an
early identification tool is necessary so that treatment
can be delivered early. The limited applicability of vari-
ous biomarkers due to the non-specific or non-sensitive
nature of these criteria, arising from the altered
physiological functions of pregnant women, could lead
to under- or over-treatment. The unnecessary use of
antibiotic therapy to treat maternal sepsis could drive
antibiotic resistance and further medical complications
[4, 19], while delay in managing sepsis can result in
many complications including organ failure, hysterectomy and death [18–20].

The use of clinical judgement is currently recom-
mended to facilitate the diagnosis of sepsis, which
should not rely solely on SSSs or laboratory tests [20].
Chorioamnionitis cases, for example, may have a clinical
diagnosis and a histological diagnosis but women might
not have a positive microbiological culture while being
treated with antibiotic for clinical chorioamnionitis [20].
Overall, the current system of diagnosing sepsis hardly
relies on the effectiveness of the early warning scores
and this might be a hindrance for validating these tools
in identifying ill patients.

In order to make SSSs more effective, addition of
biomarkers has been suggested. Procalcitonin (PCT)
shows better prediction of sepsis compared to CRP; nevertheless, there is a reported lack of accuracy
[21]. Serum PCT rises and peaks in a short time
compared to CRP and it has been found to be useful in modern clinical practice. However, there is limited
information about PCT in pregnancy, and determin-
ing the most appropriate reference range of this bio-
maker in pregnant women [22]. Its high level of
negative predicted values leads to a misleading iden-
tification of sepsis [23]. Martín and colleagues rec-
ommend the design of a scoring system based on
combined biomarker values including CRP, PCT and
lactate to aid physicians in their treatment decisions,
which has the potential to reduce the use of antibi-
otics and of culture tests [24].

With respect to compliance with SSCB, recent studies
have indicated that very few patients received all the
six interventions of the bundle within the recom-
mended 1 h and the elements were not consistently
applied [11]. This is in line with our findings. There
are possibly a number of barriers that might hinder
the timely and effective delivery of SSCB – a few
have been described such as insufficient audit and
feedback, poor teamwork and communication, inade-
quate training, lack of resources and concerns about
using SSCB in certain patients [25]. However, health
personnel also displayed positive factors in favour of
SSCB such as confidence in knowledge and skills, and
belief in the benefits of SSCB among others [25].
Since compliance with SSCB can vary among facili-
ties, regular audits, continued use, further research
into non-compliance at facility level and improving
the strengths of the staff at facility level can lead to
better implementation of SSCB.
Conclusions
In conclusion, access to a large database from maternity hospitals of a health board provided appreciable data on pregnant women with sepsis. Overall, the study indicated that mSIRS and MEWS were quite similar and weak in their characteristics in detecting sepsis in obstetric settings when compared to positive culture as a gold standard. Also, compliance with SSCB was low and administering care for all 6 elements in SSCB was not uniform. This study highlights the need for a rapid, point of care technology, incorporating a panel biomarker of greater specificity/sensitivity to support a SSS with superior prediction powers than currently available tools.

Abbreviations
AUROC: Area under the receiver operating characteristic curve; BMI: Body mass index; CRP: C-reactive protein; HDU: High dependency unit; HR: Heart rate; ICU: Intensive care unit; MEWS: Modified early warning score; MOEWS: Modified obstetric early warning score; mSIRS: Maternity systemic inflammatory response syndrome; NPV: Negative predicted value; PCT: Procalcitonin; PPV: Positive predicted value; qSOFA: Quick sepsis-related organ failure assessment; ROC: Receiver operating characteristic; RR: Respiratory rate; SBP: Systolic blood pressure; SIRS: Systemic inflammatory response syndrome; SSCB: Sepsis six care bundle; WCC: White cell count

Supplementary information
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Additional file 1: Sepsis Six Sticker.

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Authors’ contributions
Study concept and design: JG; ABM. Acquisition of data: NA; JG. Analysis and interpretation of data: NA; JG; ABM. Drafting of the manuscript: NA; ABM. Critical revision of the manuscript for important intellectual content: NA; JG; ABM. All authors (NA; JG; ABM) have read and approved the manuscript.

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Availability of data and materials
The datasets used and analysed during the current study are available from the corresponding author on reasonable request.

Declarations
Ethics approval and consent to participate
The Scientific Officer, West of Scotland Research Ethics Service reviewed the study and advised that it was considered a service evaluation that did not require formal research ethics consideration or approval. This view was endorsed by the University of Strathclyde Ethics Committee. Study design data management was approved by the local NHS Caldicott Guardian and was fully compliant with local and national clinical/data governance policies. The need for consent was deemed unnecessary according to the Strathclyde Institute of Pharmacy and Biomedical Sciences Ethics Approval committee [Reference # 0001.2016].

Consent for publication
Not required.

Competing interests
The authors report no competing interest.

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