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A step-wise approach to developing indicators to compare the performance of maternity units using hospital administrative data

Running title: How to develop maternity indicators from administrative data

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ABSTRACT

Hospital administrative data are attractive for comparing performance of maternity units due to their often large sample sizes, lack of selection bias, and the relatively low costs of accessing these data compared to conducting primary data collection. However, using administrative data to develop indicators can also present challenges including varying data quality, the limited detail on clinical risk factors and a lack of structural and user experience measures. This review illustrates how to develop performance indicators for maternity units using hospital administrative data, including methods to address the challenges administrative data pose.

Keywords: Maternity statistics, administrative data, HES, performance indicators

BACKGROUND

There is growing interest in performance monitoring and quality improvement in healthcare in the UK and elsewhere. Although quality of healthcare can be improved without measuring performance, for example, through educational programmes or clinical guidelines, accurate measurement of processes and outcomes of care is now seen as crucial for guiding service improvement.

In countries in which routinely collected hospital administrative datasets exist, these are attractive data sources for comparing performance of maternity units due to their often large sample sizes,
lack of selection bias, and the relatively low costs of accessing these data compared to conducting primary data collection. A range of performance indicators derived from these data have been proposed for both primary and secondary care. However, despite ever-high levels of public interest in the safety and quality of maternity care, many countries lack robust, easily interpretable information on even basic maternal and perinatal outcomes.

In the UK, a number of initiatives have recently been introduced, aiming to improve this situation by drawing on routinely collected clinical and hospital administrative data to measure performance and variation in care between maternity units. For example, the new National Maternity and Perinatal Audit in England, Scotland and Wales will link existing data sources, rather than introducing new bespoke data collections solely for the purposes of the audit. In Scandinavian countries, there is a long history of using medical birth registry databases, often further linked to other national databases, to conduct research into the organisation and outcomes of maternity care and to inform public health initiatives. Similar initiatives based on routine data linkage exist in Australia. A recent systematic review found that broader adoption of routine data linkage of perinatal health databases could yield substantial gains for research and surveillance. It therefore seems likely that, despite the inherent challenges of using these data, there will be a reliance on maternity indicators based, at least in part, on administrative data for some time to come.

Hospital administrative data have several advantages for describing care and outcomes. Where administrative data are readily available they are a cost-effective source of information. Where the majority of care is captured by these data, the risk of selection bias is reduced and sample sizes can be large. For example, in England >96% of all deliveries occur in NHS hospitals and are captured by administrative data (Hospital Episode Statistics, HES). Hospital administrative data also capture multiple procedures and diagnoses at the patient level, providing a rich description of patient characteristics and clinical risk factors. However, there are some important limitations of using
administrative data to develop performance indicators. There can be concerns about the accuracy and completeness of diagnosis and procedure coding,\textsuperscript{19} although there is mounting evidence that in England, most NHS trusts submit good quality data to HES.\textsuperscript{20-22} Another limitation is that not all clinical information is captured in administrative data. Some risk factors such as BMI, smoking and alcohol consumption are often not recorded. This means they cannot be taken into account in risk-adjustment for case-mix, although record linkage can extend the range of data items available and thus can improve the validity and quality of routine data. In addition, administrative data lend themselves to process and outcome indicators; measures of structural and user experience are not normally available.

As a result of these challenges and opportunities, the use of hospital administrative data for performance monitoring requires caution and a robust methodology. However, information on how to derive maternity indicators from administrative data sources is lacking. Our aim is to address this by describing a transparent approach with explicit criteria. This approach can be used by those wanting to develop performance maternity indicators using HES data, the national administrative database of the English National Health Service (NHS), or administrative data available in other healthcare settings. Furthermore, the criteria can be used by clinicians to evaluate existing performance indicators.

This approach has been used to develop indicators for the Royal College of Obstetricians and Gynaecologists’ (RCOG) Clinical Indicators Project and examples from this project will be used throughout.\textsuperscript{(10, 11)} The data source used for the examples is the HES database, containing records of admissions to NHS hospitals. Briefly, the HES database contains information on each episode of admitted patient care in the English NHS.\textsuperscript{23} Each record contains data on patient demographics (such as age, sex and ethnicity), the episode of care (e.g. hospital name, date of admission and discharge) and clinical information. Diagnoses are recorded using the International Classification of Diseases,
10th edition (ICD-10)\textsuperscript{24} and procedures using the Office of Population, Censuses and Surveys Classification of Surgical Operations and Procedures, 4th revision (OPCS-4).\textsuperscript{25} Each episode related to the delivery of a baby can also capture details about the labour and birth, such as parity, mode of delivery, gestational age and birthweight, in supplementary data fields known as the ‘maternity tail’. Each patient is assigned a unique identifier, allowing the study of longitudinal patterns of care or the number of previous births for a particular woman. For the example, delivery records were defined as those with information about a delivery in either the maternity tail or the OPCS fields in the financial year 2013/14.

DEVELOPING INDICATORS

Indicators are statistics that can describe clinical performance. The information they provide can be used for identifying possible problems and opportunities for improvement, informing policymaking, comparative benchmarking, and providing information to facilitate consumers’ choice of healthcare provider.

Building on work carried out in several surgical specialties,\textsuperscript{(20)} we assessed the suitability of using hospital administrative data for developing maternity indicators and developed a three-stage process: “identification”, “development and evaluation”, and “implementation and feedback” (summarised in Figure 1).\textsuperscript{10,11} The second stage involves evaluation against four criteria: “validity”, “statistical power”, “technical specification”, and “fairness”. Potential indicators must meet each criterion before being evaluated against the next.

Figure 1 about here

Seeking input from clinicians, methodologists and service users is a key characteristic of this indicator development process, and should be sought at each stage. Input can be via formal
consensus methods,\textsuperscript{26} or less formal round-table discussions. To minimise bias, it is important that the discussions are facilitated by a neutral chair and to ensure that all key stakeholder groups are sufficiently represented. The specific stakeholder groups that are key may vary in different settings, depending on how maternity care is organised. At the outset, all stakeholders should receive a clear brief about the aims of the initiative, the process for indicator selection and the opportunities and limitations of the data source/s. The consensus group that guided the indicator development process described in this paper consisted of obstetricians and midwives active in the English NHS, health services researchers and lay women with recent experience of English maternity care (see Acknowledgments). Lay representatives were recruited from the RCOG’s Women’s Voices Involvement Panel.\textsuperscript{27}

\textit{Stage One: Indicator identification}

To identify candidate indicators for development using administrative data, a valuable first step is a systematic review of the literature, including clinical guidelines. This also allows for an examination of associations between candidate indicators and important outcomes. Non-systematic approaches can be informative, but do not maximise the use of available evidence.\textsuperscript{8} In addition to identifying indicators from the literature, suggestions can be sought from stakeholders with an interest in measuring the performance of maternity services via surveys or face-to-face meetings, thereby reducing the potential impact of publication bias.

To provide a broad understanding of the performance and quality of a healthcare service as a whole it is important that a suite of indicators is “balanced”. A balanced suite would ideally include indicators relating to the structure of care, the processes of care, or the outcomes of the care received\textsuperscript{6} throughout the care pathway, and including measures of user experience. Indicators derived from administrative data will tend to focus on process and outcome indicators as structural and user experience measures are not normally available in these datasets. However, they are
nonetheless important for understanding many care outcomes. Balancing the types of measures used to evaluate performance can also help to minimise the risk of indicators being taken out of context and in that way misinforming quality improvement initiatives.

**Stage Two: Indicator development and evaluation**

Given the challenges of using administrative data it is important to rigorously evaluate candidate indicators to address these issues as far as possible. In our process this entailed evaluation against four criteria in turn: “validity”, “statistical power”, “technical specification”, and “fairness”.

**Criterion One: Validity**

Clinical and lay input should be sought to identify which of the identified indicators are considered to be clinically meaningful, or in other words, to measure aspects of the service or the quality of care provided that are relevant to patients. For an indicator to be considered valid, it must also be likely that a difference in the indicator reflects a difference the quality of care, and a specific direction should reflect better quality. For example, a higher rate of “obstetric anal sphincter injury” can be thought to reflect poorer obstetric care. Indicators not meeting this criterion should be dropped at this stage. Examples of decisions to include, refine and exclude indicators based on assessments of validity in the RCOG Maternity Indicators Project are provided in Appendix S1.

A key consideration when using hospital administrative data to develop indicators of quality of care is whether denominators and numerators can be adequately captured. Once an indicator was identified as valid, input from clinicians was used to define the appropriate “denominator” (the group of patients for whom the indicator is relevant) and “numerator” (the state or the event of which the frequency is captured by the indicator). For example, an indicator reflecting the use of elective CS before 39 weeks without clinical indication would have as its denominator the number of patients who had an elective CS without a recorded clinical indication (e.g. gestational hypertension,

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gestational diabetes, or poor fetal growth) and as its numerator the number of patients in this group who had an elective CS before 39 weeks.\textsuperscript{11}

Not all variation in performance indicators will reflect variation in quality of care. Factors including random fluctuations, differences in data quality, and the case-mix of patients between hospitals may account for a large part of the observed variation. Conclusions about quality of care can only be reasonably drawn after differences due to these other factors are excluded. With criteria two, three and four we describe a transparent approach to address these issues in administrative datasets.

\textit{Criterion Two: Statistical power}

An advantage of using administrative data to develop indicators is their often large sample size. However, even in large datasets, where an event or a procedure that forms part of an indicator is rare, the statistical power to identify providers with truly poor performance is low. In this situation, no evidence of poor performance cannot be taken as evidence of acceptable performance.\textsuperscript{29}

Indicators should be reported at a level (clinic, hospital, NHS trust) which is appropriate to how care is commissioned and provided. However, where numbers within a unit are too small, a higher-level unit of analysis, or a longer timeframe should be considered. If this is not appropriate (given the way that care is commissioned or provided) the indicator cannot be judged to have met this statistical power criterion. We rejected some maternity indicators due to a small number of events per hospital per year. For example, the maternal mortality rate in the UK is 8.5/100,000 pregnancies.\textsuperscript{30}

Therefore, the ‘signal to noise’ ratio for this measure is too low to detect true differences between hospitals.\textsuperscript{31,32} In this situation, composite indicators may be appropriate. For example, a composite maternal morbidity indicator has been proposed using Australian routine hospital data.\textsuperscript{33,34}
Criterion Three: Feasibility of technical specification

As administrative data are collected for administrative purposes rather than for research or quality improvement, not all data items required for specific indicators may be adequately captured. Valid and adequately powered indicators should therefore next be evaluated in terms of their technical specification. This comprises a detailed assessment of the available data source(s) to establish how well patient populations, important case-mix differences and the procedures or outcomes that define the indicator can be captured. In HES, this involves exploring the diagnosis (ICD-10) and procedure (OPCS-4) codes which can be used to define the indicator in preliminary analyses. The technical specification of the inclusions and exclusions defined in step 1 should also be evaluated.

Where data required to construct the indicators and identify the units of analysis are available, an assessment of the data quality and completeness should also be conducted. Identifying data quality issues that would affect our ability to define the appropriate populations for each indicator allowed us to be confident that indicators were based on data that met minimum standards. We propose assessing data quality overall, and by hospital, using three main methods:11

- Investigation of the proportion of missing data
- Internal consistency between data items within HES21,35. For example, we excluded hospitals in which less than 90% of the records had consistency of mode of delivery between the main HES record and the maternity tail (Appendix S1).
- Comparison with results from external studies

Examples of data quality assessments conducted as part of the Maternity Indicators Project are provided in Appendix S2. Full details of how data quality was assessed have been published elsewhere.11
Another example of evaluating the technical specification of an indicator is a recent study that explored whether a composite maternal morbidity indicator developed using Australian routine data could be derived from HES data. This study found that the quality of the relevant HES data meant that 11 conditions that were included in the Australian indicator would have to be excluded from the English indicator. These included eclampsia, obstetric embolism and cardiac arrest/failure, which are associated with increased risk of maternal mortality in the UK, making the resultant indicator questionable in its ability to accurately estimate maternal morbidity during childbirth in England.

*Criterion Four: Fairness*

Patient characteristics may influence indications for procedures and treatments, as well as influencing outcomes. Indicators should only be used for comparative purposes where adequate adjustment has been made for key case-mix differences between populations of patients. Calculating indicators without appropriate risk-adjustment may give rise to misleading results. A number of questions should be evaluated as part of robust risk-adjustment (Box 2).

Hospital administrative data are able to capture multiple procedures and diagnoses at an individual level, providing a rich description of the case-mix of patient characteristics and clinical risk factors. However, not all clinical information is captured; risk factors such as BMI, smoking and alcohol consumption are not recorded, meaning they cannot be accounted for.

In maternity care, certain pregnancy characteristics can have a large impact on the care provided and on outcomes. To ensure that the indicators would allow fair comparisons among hospitals, we decided to focus on women with singleton, term, cephalic deliveries, whose maternity care is most affected by between-hospital and provider variation in clinical practices. Multiple births, preterm births and breech deliveries require very different management. Remaining differences in case-mix between hospitals can be addressed in several ways. First, indicators can be stratified by a
clinical condition that has a major influence on outcomes. For example, we stratified maternity indicators by parity (nulliparous and multiparous). Second, risk-adjustment using a regression model can be used. Further information on the methods and impact of risk adjustment in the RCOG’s maternity indicators project is provided in Appendix S3. Identifying which factors to include in these risk adjustment models is complex, requiring knowledge of the relevant factors, statistical expertise and adequate data.

Following evaluation against these four criteria of: “validity”, “statistical power”, “technical specification”, and “fairness” four in turn, 18 maternity indicators were developed from HES data (Table 1).

Table 1 about here

Stage Three: Implementation and feedback

Given high levels of public interest in the quality of maternity care, and the challenges associated with using administrative data for this purpose, the use of indicators for performance assessment derived from administrative data needs to be cautiously implemented. A feasibility phase in which hospital-specific results are published anonymously can generate buy-in from those who will ultimately use the indicators. Individual, personalised feedback of results to the hospitals may also give them an opportunity to address identified data quality issues. Also, careful consideration should be given to the best methods for reporting results to achieve maximum impact with the intended audiences. This may be static reports, interactive online formats, and/or a series of local or regional discussion meetings. Finally, it is important to encourage those who use indicators not to interpret the results of individual indicators in isolation, but to look at a suite as a whole, considering possible relationships between indicators. For example, in maternity care there is an association between lower pre-labour CS rates and therefore higher vaginal delivery rates on the
one hand but also higher emergency CS rates, which in turn can influence outcomes such as length of stay or readmission post-delivery.

CONCLUDING REMARKS

There is international interest in using indicators derived from administrative data to drive improvements in maternity care but information on how to derive indicators from administrative data, addressing the challenges presented by these data, is lacking. We present an approach for developing indicators using administrative data that has been well received by healthcare professionals and addresses many of the challenges of using administrative data for this purpose. Key features of this process are explicit data quality checks, risk-adjusting hospital results for differences in patient case-mix, and clinical and lay input at all stages. Some of the indicators developed have already been incorporated into local monitoring systems and national outcome frameworks (Box 1) and will be developed further as part of the new National Maternity and Perinatal Audit. Our indicator development process is also more “streamlined” than others, with three steps and four evaluation criteria. This supportive approach also included collaboration between those developing indicators and those whose performance they are designed to monitor.

Box 1 about here

Overall, hospital administrative data sources are attractive for comparing performance of maternity units within and between countries due to their often large sample sizes, lack of selection bias, and the relatively low costs of accessing these data compared to those of conducting primary data collection. However, using administrative data to develop maternity indicators also present challenges, and the development of the indicators described in this review has triggered debate about the use of administrative data for this purpose. Such debate has included concerns about the accuracy and completeness of coding, that data quality may vary between healthcare providers, the
lack of detail on risk factors such as BMI, smoking and alcohol consumption and the absence of structural and user experience measures. However, a systematic review of discharge coding accuracy in administrative UK data found that primary diagnosis accuracy improved from 73.8% to 96.0% in the last decade, concluding that administrative data are sufficiently robust to use for research and managerial decision-making. For each indicator proposed, careful evaluation of how well it can be derived from administrative data to meet “validity”, “statistical power”, “technical specification” and “fairness criteria” allows some of these challenges to be addressed. This transparent approach to developing indicators using administrative data could also be applied in other specialties, for primary care, and for international, national or regional comparisons.

Hospital administrative data are not the perfect data source for developing indicators of maternity care quality. Because administrative data are not collected for research purposes the performance indicators available may differ from core outcome sets used in clinical trials that focus on quality improvement and safety. The development of more clinically detailed routine maternity datasets will ultimately allow for improvement of existing indicators, and the development of new indicators, producing a more balanced picture of the quality of maternity care. However, until centrally-available electronic maternity records become the norm, routine hospital administrative data, linked with other sources of clinical and user experience data where possible, will be the key data source for performance indicators. Some countries, such as Denmark, Finland, Norway and Sweden, are ahead of the game in integrating data linkage into their routine perinatal health surveillance systems and making these data available for research, but this is not a universal practice even in high-income countries with access to electronic hospital administrative data. Standardisation of performance measures derived from administrative data research would be desirable to facilitate comparisons both nationally and internationally.
It seems likely that there will be a reliance on maternity indicators based, at least in part, on administrative data for some time to come. In light of this, methods for addressing the challenges posed by administrative data for the development of performance indicators are sorely needed. The transparent approach detailed in this paper aims to contribute to this effort. Our approach has led to the development of maternity indicators that have been adopted at a local and national-level, and addresses many of the issues raised about the usefulness of administrative data for performance monitoring.

**Box 1: examples of use of indicator definitions or data**

Information from the maternity indicators project has already been used by trusts and incorporated into local monitoring systems, national outcomes frameworks. For example:

- The definition of the indicator “Elective caesarean section without indication before 39 weeks of gestation” has been proposed for the Clinical Commissioning Group Outcomes Indicator Set.
- Several Clinical Networks held regional workshops to encourage trusts within the same region to compare results and reflect on the cause of any differences in practices or outcomes.
- A number of indicators from this project have been included in regional dashboards.
- The indicators have been used by trusts in the following ways (based on a small evaluation survey carried out in May 2016; n=19 trusts):
  - Discussed with clinical board/Senior Management Team: 55% (n=11)
  - Led to an internal audit: 30% (n=6)
  - Led to an investigation of data collection/coding/provision: 45% (n=9)
  - Led to a change in data systems/ability to provide data in the future: 25% (n=5)
Case Mix
How big are the important case-mix differences between hospitals/trusts?

Data
Does sufficient detail on case-mix exist in the available data?
If not, could data linkage be used to obtain these data from other sources?

Unmeasured Confounding
Which factors do not have data available which could result in unmeasured confounding?

Impact of Adjustment
What is the impact of risk-adjustment on the differences between the hospitals/trusts?

Box 2: Risk Adjustment: Questions to Ask

Ethical approval
The study is exempt from UK National Research Ethics Service (NRES) approval because it involved the analysis of an existing dataset of anonymized data for service evaluation. HES data were made available by NHS Digital (Copyright 2015, Re-used with the permission of NHS Digital. All rights reserved.) Approvals for the use of anonymized HES data were obtained as part of the standard NHS Digital data access process.

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 Contributors
RSG, HEK, IGU, EM and JvdM conceived the review. FC, HEK and IGU performed the data analysis for the Clinical Indicators Project that informed this review, with technical support from DAC and advice from JvdM and EM. All authors contributed to the writing of the manuscript.
Disclosure of interests

The authors declare that they have no conflicts of interest. The ICMJE disclosure forms are available as online supporting information.

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References

31. Mant J, Hicks N. Detecting differences in quality of care: the sensitivity of measures of process
Table 1: Indicators developed from HES for the RCOG’s Maternity Indicators Project

<table>
<thead>
<tr>
<th>Indicator Description</th>
<th>Population subset</th>
</tr>
</thead>
<tbody>
<tr>
<td>1. UNASSISTED VAGINAL DELIVERIES</td>
<td></td>
</tr>
<tr>
<td>1a) Proportion of spontaneous, unassisted vaginal deliveries</td>
<td>Primip/Multip</td>
</tr>
<tr>
<td>2. INDICATORS RELATED TO INDUCTION OF LABOUR</td>
<td></td>
</tr>
<tr>
<td>2a) Proportion of induced labours</td>
<td>Primip/Multip</td>
</tr>
<tr>
<td>2b) Proportion of induced labours in deliveries between 37 and 39 weeks of gestation</td>
<td>Primip/Multip</td>
</tr>
<tr>
<td>2c) Proportion of induced labours in deliveries ≥42 weeks of gestation</td>
<td>Primip/Multip</td>
</tr>
<tr>
<td>3. INDICATORS RELATING TO CAESAREAN SECTION</td>
<td></td>
</tr>
<tr>
<td>3a) Proportion of deliveries by caesarean section</td>
<td>Primip/Multip</td>
</tr>
<tr>
<td>3b) Proportion of induced labours resulting in emergency caesarean section</td>
<td>Primip/Multip</td>
</tr>
<tr>
<td>3c) Proportion of spontaneous labours resulting emergency caesarean section</td>
<td>Primip/Multip</td>
</tr>
<tr>
<td>3d) Proportion of prelabour caesarean sections</td>
<td>Primip/Multip</td>
</tr>
<tr>
<td>3e) Proportion of prelabour caesarean sections performed before 39 weeks of gestation</td>
<td>Pre</td>
</tr>
<tr>
<td>3f) Proportion of vaginal births following a primary caesarean section (VBAC)</td>
<td>Multip</td>
</tr>
<tr>
<td>4. INVOLVEMENT OF INSTRUMENTS</td>
<td></td>
</tr>
<tr>
<td>4a) Proportion of deliveries involving instruments</td>
<td>Primip/Multip</td>
</tr>
<tr>
<td>5. EPISIOTOMY</td>
<td></td>
</tr>
<tr>
<td>5a) Proportion of episiotomies among vaginal deliveries</td>
<td>Primip/Multip</td>
</tr>
<tr>
<td>5b) Proportion of episiotomies among instrumental deliveries</td>
<td>F/Va</td>
</tr>
<tr>
<td>6. INDICATORS RELATING TO 3rd AND 4th DEGREE TEARS</td>
<td></td>
</tr>
<tr>
<td>6a) Proportion of third- and fourth-degree perineal tears among vaginal deliveries</td>
<td>Primip/Multip</td>
</tr>
<tr>
<td>6b) Proportion of third- and fourth-degree perineal tears among unassisted vaginal</td>
<td>Primip/Multip</td>
</tr>
<tr>
<td>deliveries</td>
<td></td>
</tr>
<tr>
<td>6c) Proportion of third- and fourth-degree perineal tears among assisted vaginal</td>
<td>Primip/Multip</td>
</tr>
<tr>
<td>deliveries</td>
<td></td>
</tr>
<tr>
<td>7. ADMISSIONS TO HOSPITAL FOLLOWING DELIVERY</td>
<td></td>
</tr>
<tr>
<td>7a) Unplanned maternal readmission to hospital within 42 days of delivery</td>
<td>V/CS</td>
</tr>
<tr>
<td>7b) Unplanned neonatal readmission to hospital within 28 days of birth</td>
<td>NB</td>
</tr>
</tbody>
</table>

Footnote: For all indicators, multiple and preterm deliveries were excluded. Women who delivered a baby with a non-cephalic presentation were also excluded, apart from for indicators 3e and 7b. Primip=Primiparous; Multip=Multiparous; CS=Caesarean section deliveries; F=Forceps, NB= Normal Birthweight Infants; Pre= Subset of prelabour caesarean section deliveries including women with non-cephalic presentation OR where 1 or 2 previous caesarean sections; Va=Vacuum
HOW TO DEVELOP CLINICAL INDICATORS

1: IDENTIFICATION
Identify candidate indicators with a systematic literature review
Use clinical and lay input to decide which are important

2: DEVELOPMENT AND EVALUATION
Evaluate indicators in terms of validity, statistical power, technical specification and fairness in turn.
If an indicator fails to meet one criteria, remove it

2A: VALIDITY
Are differences in an indicator likely to reflect quality of care?

2B: STATISTICAL POWER
What is the chance of detecting a true outlier?

2C: TECHNICAL SPEC
How well can the data define patients, procedures or outcomes for an indicator?

2D: FAIRNESS
How different are patients treated by different units? Can we capture & adjust for this?

3) IMPLEMENTATION AND FEEDBACK
- Engage users early
- Publish indicators anonymously first to build trust and drive up data quality
- Interpret related indicators together, not alone

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