# Parametric Collapsibility and the Lack of Moderating Effects in Contingency Tables with a Dichotomous Response Variable 

By NANNY WERMUTH $\dagger$<br>University of Mainz, Federal Republic of Germany

[Received May 1986. Final revision May 1987]
SUMMARY
We consider contingency tables having one variable specified as a response with just two categories. We look at conditions for collapsibility of a symmetric and a directed measure of association, the odds-ratio and the relative risk: situations are discussed under which equal partial associations coincide with the corresponding marginal association. Contrary to the odds-ratio the relative risk is collapsible, if there are independencies in the marginal distribution of the influencing variables. This fact is exploited to derive conditions for the lack of a moderating effect, the latter being a much discussed concept in the social science literature.

Keywords: INDEPENDENT REGRESSORS; ODDS-RATIO; RELATIVE RISK; YULE-SIMPSON PARADOX

## 1. INTRODUCTION

In many empirical studies, one wants to investigate how a response variable with just two categories depends on several qualitative variables. Data from such studies can be summarised in contingency tables containing at least one dichotomous variable, the response. Typically, there are some variables for which potential influences are of main interest and some others that play the role of only background variables: one needs to include the latter explicitly in an analysis, because they might modify one of the associations of interest, but one hopes to be able to report the results irrespective of the categories of the background variables. Typical examples are simple multicentre clinical trials, in which the categories of the response variable $\left(V_{1}\right)$ are success or failure of a treatment, those of the background variable $\left(V_{3}\right)$ are the different clinics or sites and those of the main variable of interest $\left(V_{2}\right)$ are the treatment types. Reporting results in such studies irrespective of site can be justified if one has strongly consistent results within sites and these agree with the result overall, that is, for the data combined from all sites. This is a description by example, in which the so-called moderating effect of a background variable is lacking. The concept of moderator variables is much discussed in the social science literature. However, its implementation has remained unsatisfactory. For a survey see Zedeck (1971).

Moderator variables are, in the social sciences, mainly of interest in non-experimental situations like the following. The risk that a person becomes depressive is known to increase with the occurrence of life events like birth, death, severe illness or separation of friends. Since it is, in general, not possible to control such events, it becomes important to understand conditions which are likely to change or moderate this known relationship. Thus, one wants to determine which of the personality characteristics, environmental conditions, coping strategies or therapies are important moderator variables. One wants to estimate moderating effects and to understand under which conditions such effects cannot occur.

Associated statistical tasks are to assess the relative importance of an additional influencing or regressor variable, to estimate the bias introduced by neglecting an important influencing variable and to state conditions under which a measure of association is unaffected by adding

[^0]a variable to an analysis. All of these aspects are well understood in the case of exclusively linear dependencies (Goldberger, 1964, chapter 10). For symmetric associations among discrete variables the last aspect has been called (parametric) collapsibility and was studied by Yule (1900), Simpson (1951), Bishop (1971), Whittemore (1978) and Ekholm (1985). In this paper we treat the situation in which all variables under study are discrete and there is a dichotomous response variable.

There are two basic aspects to the lack of a moderating effect: (1) associations have to coincide for all levels of the moderator variable and (2) the conclusions regarding the associations have to remain unchanged if the moderator variable is excluded from the analysis. Correspondingly, we speak of strongly consistent results in terms of a given measure of association, if all partial associations within sites are equal, and we say that a measure of association is collapsible if also the equal partial associations coincide with the marginal association. We discuss in section 2 necessary and sufficient conditions for the collapsibility of the odds-ratio and the relative risk in the simplest case of a $2^{3}$-contingency table. In section 3 we give necessary and sufficient conditions for the lack of a moderating effect in those $2^{3}$-tables, in which equal partial relative risks are always collapsible. Nontrivial sufficient conditions for the lack of a moderating effect on the relative risk are derived in section 4 for general tables with nonzero cell probabilities. Finally, we show in section 5 how, for data in a $2 \times 3 \times 3 \times 2$ table, where the response depends on all three of the remaining variables, the hypothesis of no moderating effect on the relative risk fits the data well.

## 2. CONDITIONS FOR COLLAPSIBILITY OF ODDS-RATIOS AND OF RELATIVE RISKS

For a $2 \times J \times K$ table of probabilities we use the indices $i, j$, and $k$ for the categories of the variables $V_{1}, V_{2}$, and $V_{3}$, respectively: denote by $\Pi_{i j k}$ the joint probability $P\left(V_{1}=i, V_{2}=j\right.$, $\left.V_{3}=k\right)>0$. To make the interpretations of different measures of association more vivid, we name the variables as in simple multicentre studies: $V_{1}$ as outcome or response, $V_{2}$ as treatment and $V_{3}$ as clinic or site. For our discussion on collapsibility we need marginal probabilities, such as

$$
\Pi_{i j+}=\Sigma_{k} \Pi_{i j k}=P\left(V_{1}=i, V_{2}=j\right)
$$

which are called joint marginal probabilities in the $2 \times J$ marginal contingency table. Furthermore, we use conditional probabilities such as, for $V_{1}=1$ given $V_{2}=j$ and $V_{3}=k$,

$$
\Pi_{1 \mid j k}=\Pi_{1 j k} / \Pi_{+j k} .
$$

In the context of log-linear models for contingency tables (Birch, 1963) the natural measures of association are logarithms of so-called odds-ratios. For instance, the odds-ratios for outcome and treatments 1 and 2 within sites are

$$
\begin{equation*}
\frac{\Pi_{11 k} \Pi_{22 k}}{\Pi_{12 k} \Pi_{21 k}}, \quad k=1, \ldots, K \tag{2.1}
\end{equation*}
$$

They are also called partial odds-ratios for $V_{1}$ and $V_{2}$ given $V_{3}$.
In epidemiological studies, the measure of association used before the advent of log-linear models was the relative risk (see, e.g., Hill, 1962). For instance, the relative risks for $V_{1}=1$ under treatment 1 compared to treatment 2 within sites are the following ratios of conditional probabilities:

$$
\begin{equation*}
\frac{\Pi_{1 \mid 1 k}}{\Pi_{1 \mid 2 k}}, \quad k=1, \ldots, K . \tag{2.2}
\end{equation*}
$$

They are also labelled the partial relative risks for $V_{1}=1$, given $V_{3}$.
One obtains the corresponding marginal measures of association from the probabilities in the marginal table of $V_{1}$ and $V_{2}$, or, to put it differently, after replacing " $k$ " in (2.1) and (2.2) by
" + ". Results are said to be weakly consistent within sites if the partial associations given sites lead to similar interpretations regarding the success ( $V_{1}=1$ ) of treatment $1\left(V_{2}=1\right)$ in comparison to the success of treatment 2: treatment 1 appears to be better, within sites, than treatment 2 if the odds-ratios in (2.1) or the relative risks in (2.2) are larger than one. Situations in which weakly consistent results appear reversed overall have been named the Yule-Simpson paradox (Yule 1900; Simpson 1951).

If, given treatment 1 , response is independent of sites then we speak of conditional independence of $V_{1}$ and $V_{3}$ given $V_{2}=1$ and write $V_{1} \Perp V_{3} \mid\left(V_{2}=1\right)$. In that case one of the following equivalent conditions, stated here without proof, holds:
(iii)

$$
\begin{align*}
\Pi_{1 \mid 11} & =\Pi_{1 \mid 1+}  \tag{i}\\
\frac{\Pi_{111} \Pi_{212}}{\Pi_{112} \Pi_{211}} & =1 \tag{ii}
\end{align*}
$$

$$
\begin{equation*}
\frac{\Pi_{1 \mid 11}}{\Pi_{1 \mid 12}}=1 . \tag{2.3}
\end{equation*}
$$

For $V_{1} \Perp V_{3} \mid\left(V_{2}=1\right)$ and $V_{1} \Perp V_{3} \mid\left(V_{2}=2\right)$ we write, in brief, $V_{1} \Perp V_{3} \mid V_{2}$ and speak of conditional independence of $V_{1}$ and $V_{3}$ given $V_{2}$.

The more standard description for $V_{1} \Perp V_{3} \mid V_{2}$ is $\Pi_{i j k}=\Pi_{i j+} \Pi_{+j k} / \Pi_{+j+}$, and, for $V_{3} \Perp\left(V_{1}\right.$, $V_{2}$ ), the complete independence of $V_{3}$ from $V_{2}$ and $V_{1}$, taken jointly, it is $\Pi_{i j k}=\Pi_{i j+} \Pi_{++k}$, where both hold for all $(i, j, k)$. It is known (see, e.g., Birch, 1963) that $V_{3} \Perp\left(V_{1}, V_{2}\right)$ is equivalent to $V_{1} \Perp V_{3} \mid V_{2}$ and $V_{2} \Perp V_{3} \mid V_{1}$. We now turn to conditions for collapsibility.

Proposition 1. In $2^{3}$-contingency tables necessary and sufficient conditions for results, which are strongly consistent, also to be collapsible are
(i) for odds-ratios of $V_{1}$ and $V_{2}$ (2.1): $V_{1} \Perp V_{3} \mid V_{2}$ or $V_{2} \Perp V_{3} \mid V_{1}$,
(ii) for relative risks of $V_{1}(2.2): V_{1} \Perp V_{3} \mid V_{2}$ or $V_{2} \Perp V_{3}$.

Statement (i) is an immediate consequence of the conditions for collapsibility of log-linear interaction parmeters (Whittemore, 1978), and (ii) is proven in the Appendix.

Remark 1. As far as collapsibility is concerned, odds-ratios behave analogously to concentrations $\sigma^{i j}$. These are elements in the inverse of a positive definite covariance matrix $\Sigma$ and they are simple multiples of partial correlation coefficients given all other variables (see, e.g., Wermuth, 1976). For trivariate normal distributions the following three statements are equivalent:
(i) $V_{i} \Perp V_{j} \mid V_{k}$,
(ii) $\sigma^{i j}=0$,
(iii) $\rho_{i j, k}=0$,
where $i, j$ and $k$ are distinct elements of $\{1,2,3\}$ and $\rho_{i j, k}$ denotes a partial correlation coefficient. The marginal concentration between $V_{1}$ and $V_{2}, \sigma^{12.3}$, is known to be related to the partial one, $\sigma^{12}$ (see, e.g., Wermuth, 1980) through

$$
\sigma^{12.3}=\sigma^{12}-\sigma^{13} \sigma^{23} / \sigma^{33}
$$

so that, from $\sigma^{33}>0$, by positive definiteness, the claimed analogy becomes evident. This is not surprising, if one considers the exponential family parametrisations (see, e.g., Dempster, 1971) of a joint normal and of a multinomial distribution for three variables. Concentrations are there the natural parameters for the former and log-odds ratios for the latter, provided the prerequisite for collapsibility is satisfied, that is, provided one has equal partial odds-ratios.

Remark 2. Relative risks behave similarly, but not completely analogously, to regression coefficients in bivariate normal regressions. Let $\beta_{12.3}$ denote the partial regression coefficient, the coefficient of $V_{2}$ in a linear regression of $V_{1}$ on $V_{2}$ and $V_{3}$, and $\beta_{12}$, the marginal regression coefficient, the coefficient of $V_{2}$ in a linear regression of $V_{1}$ on $V_{2}$ alone. Then it is known (Currie and Korabinski, 1984) that the two are related through

$$
\beta_{12.3}=\beta_{12} \frac{\left(\rho_{12}-\rho_{13} \rho_{23}\right)}{\rho_{12}\left(1-\rho_{23}^{2}\right)}
$$

where the $\rho$ 's are marginal correlation coefficients. This makes it plain that the regression coefficient is collapsible in the case of $\beta_{12.3} \neq 0\left(\right.$ not $\left.V_{1} \Perp V_{2} \mid V_{3}\right)$ if and only if

$$
\rho_{23}=0 \quad \text { or } \quad \rho_{13}=\rho_{12} \rho_{23}
$$

which for joint normal distributions is equivalent to requiring that

$$
V_{2} \Perp V_{3} \quad \text { or } \quad V_{1} \Perp V_{3} \mid V_{2} .
$$

The difference, which remains to relative risks, concerns the strong consistency in terms of the analogous measures: concentrations and odds-ratios. In normal distribution theory the conditional associations between $V_{1}$ and $V_{2}$, given different levels of $V_{3}$, are equal by definition, which is reflected in just one possible value for the partial concentration $\sigma^{12}$ (or the partial correlation coefficient $\rho_{12.3}$ ). The relative risk can however, for $V_{2} \Perp V_{3}$, be collapsible even if one has only weakly consistent results in terms of odds-ratios. An example is given in Table 1.

TABLE 1
Example for collapsible relative risks* in spite of unequal partial odds-ratios

|  | $k=1$ |  | $k=2$ |  | Sum over $k$ |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | $j=1$ | $j=2$ | $j=1$ | $j=2$ | $j=1$ | $j=2$ |
| $i=1$ | $\begin{array}{r} 6 \\ (30 \%) \end{array}$ | $\begin{array}{r} 40 \\ (20 \%) \end{array}$ | $\begin{array}{r} 30 \\ (75 \%) \end{array}$ | $\begin{array}{r} 200 \\ (50 \%) \end{array}$ | $\begin{array}{r} 36 \\ (60 \%) \end{array}$ | $\begin{array}{r} 240 \\ (40 \%) \end{array}$ |
| $i=2$ | 14 | 160 | 10 | 200 | 24 | 360 |
| Sum | 20 | 200 | 40 | 400 | 60 | 600 |
|  | $\frac{\Pi_{1 \mid 11}}{\Pi_{1 \mid 21}}=1.5$ |  | $\Pi_{1[12}=1.5$ |  | $\Pi_{1!1+}{ }_{-} 1.5$ |  |

*All probabilities $\Pi_{i j k}$ are multiplied by 660 .

Remark 3. If outcome is independent of treatment, given sites $\left(V_{1} \Perp V_{2} \mid V_{3}\right.$ ), then the results are strongly consistent in terms of relative risks and in terms of odds-ratios, and either both or none of these measures of associations are collapsible. Examples for the latter case are special versions of the Yule-Simpson paradox described for instance by Birch (1963, (5.1)) or by Bishop, Fienberg and Holland (1975, p. 41). To see the former, note that $V_{1} \Perp V_{2} \mid V_{3}$ and $V_{2} \Perp V_{3} \mid V_{1}$ imply $V_{2} \Perp\left(V_{3}, V_{1}\right)$ and hence $V_{2} \Perp V_{3}$. Similarly, $V_{1} \Perp V_{2} \mid V_{3}$ and $V_{2} \Perp V_{3}$ imply $V_{2} \mathbb{H}\left(V_{1}, V_{3}\right)$, and in particular $V_{2} \Perp V_{3} \mid V_{1}$.

Remark 4. If outcome depends conditionally on both of treatments and sites, then at most
one of the measures of association is collapsible. To see this, note first that, for $V_{1} \Perp V_{2} \mid V_{3}$ not to be satisfied, strongly consistent results in terms of odds-ratios and relative risks imply $V_{1} \Perp V_{3} \mid V_{2}$ (for details see e.g., Wermuth, 1986) and hence contradict an assumption. Note secondly that the two conditions for collapsibility of odds-ratios and relative risks cannot both be satisfied in tables with neither $V_{1} \Perp V_{2} \mid V_{3}$ nor $V_{1} \Perp V_{3} \mid V_{2}$ : Birch (1963) has shown that in $2 \times J \times K$ tables the two independencies $V_{2} \Perp V_{3} \mid V_{1}$ and $V_{2} \Perp V_{3}$ together imply that one of the stronger hypotheses $V_{2} \Perp\left(V_{1}, V_{3}\right)$ or $V_{3} \Perp\left(V_{1}, V_{2}\right)$ has to hold. The claim then follows, because $V_{1} \Perp V_{2} \mid V_{3}$ and $V_{1} \Perp V_{3} \mid V_{2}$ are necessary conditions for $V_{2} \Perp\left(V_{1}, V_{3}\right)$ and for $V_{3} \Perp\left(V_{1}, V_{2}\right)$, respectively.

One conclusion, for data analyses, from Proposition 1 and the example in Table 1 is that collapsibility of relative risks offers new possibilities for giving more condensed descriptions of results in situations where the partial odds-ratios do not coincide. We therefore investigate tables satisfying $V_{2} \Perp V_{3}$ next.

## 3. EFFECTS OF ONE MARGINAL INDEPENDENCE IN A $2^{3}$-TABLE

A $2^{3}$-table with $V_{2} \Perp V_{3}$ can be viewed as the simplest situation in which a dichotomous response variable $\left(V_{1}\right)$ has two qualitative variables as possible influences and these are independent of each other. For $V_{2} \Perp V_{3}$ one has, by definition, for $j=1,2$ and $k=1,2$,

$$
\begin{equation*}
\Pi_{+j k} / \Pi_{+j+}=\Pi_{++k}, \tag{i}
\end{equation*}
$$

or

$$
\begin{equation*}
\frac{\Pi_{+11} \Pi_{+22}}{\Pi_{+21} \Pi_{+12}}=1 \tag{ii}
\end{equation*}
$$

or
(iii)

$$
\Pi_{+1 k} / \Pi_{+1+}=\Pi_{+2 k} \Pi_{+2+}
$$

This restriction on the probabilities in the marginal contingency table of $V_{2}$ and $V_{3}$ has effects on the possible associations in the joint table, to see which, we note first that then

$$
\begin{equation*}
\frac{\Pi_{1 \mid 1+}}{\Pi_{1 \mid 2+}}=a \frac{\Pi_{1 \mid 11}}{\Pi_{1 \mid 21}}+(1-a) \frac{\Pi_{1 \mid 12}}{\Pi_{1 \mid 22}} \tag{3.2}
\end{equation*}
$$

with $a=\Pi_{1 \mid 21} \Pi_{++1} /\left(\Pi_{1 \mid 21} \Pi_{++1}+\Pi_{1 \mid 22} \Pi_{++2}\right)$.
Thus, contrary to the case for general $2^{3}$-tables, the marginal relative risk is a weighted average of the corresponding partial quantities, with positive weights adding to one. The implications of (3.2) are (1) that strongly consistent results in terms of relative risks are always collapsible, i.e., in agreement with the result overall, and (2) that weakly consistent results within sites can never appear reversed overall, that is, the Yule-Simpson paradox cannot occur. The last statement follows after noting that, in tables with a dichotomous response, weak consistency in terms of one measure of association implies weak consistency in terms of the other.

Proposition 2: In $2^{3}$-contingency tables with $V_{2} \Perp V_{3}$ the necessary and sufficient condition for the lack of a moderating effect of $V_{3}$ is
(i) for odds-ratios of $V_{1}$ and $V_{2}: V_{2} \Perp V_{3} \mid V_{1}$,
(ii) for relative risks of $V_{1}=1: V_{2} \Perp V_{3} \mid\left(V_{1}=1\right)$.

A moderating effect is lacking, if one has equal partial associations and these are collapsible. Since $V_{2} \Perp V_{3}$ is sufficient for collapsibility of equal relative risks, Proposition 2 (ii) results by
noting the following equivalent statements for $2^{3}$-tables with $V_{2} \Perp V_{3}$ :
(i)

$$
\begin{equation*}
\frac{\Pi_{1 \mid 11}}{\Pi_{1 \mid 21}}=\frac{\Pi_{1 \mid 12}}{\Pi_{1 \mid 22}} \tag{3.3}
\end{equation*}
$$

$$
\begin{equation*}
\frac{\Pi_{111} \Pi_{122}}{\Pi_{121} \Pi_{112}}=1 \tag{ii}
\end{equation*}
$$

(iii)

$$
V_{2} \Perp V_{3} \mid\left(V_{1}=1\right) .
$$

For the proof of Proposition 2 (i) note first from Proposition 1 that conditions for the lack of a moderating effect on the odds-ratio are included in those for the collapsibility: if $V_{1} \Perp V_{3} \mid V_{2}$ or $V_{2} \Perp V_{3} \mid V_{1}$ then one has equal partial odds-ratios. Note secondly that these conditions are still necessary and sufficient in $2^{3}$-tables with $V_{2} \mathbb{\Perp} V_{3}$, but that conditional independencies of variable pairs have different interpretations,
(i) $\quad V_{1} \Perp V_{2} \mid V_{3}$ if and only if $V_{2} \Perp\left(V_{1}, V_{3}\right)$,
(ii) $\quad V_{1} \Perp V_{3} \mid V_{2}$ if and only if $V_{3} \Perp\left(V_{1}, V_{2}\right)$,
(iii) $\quad V_{2} \Perp V_{3} \mid V_{1}$ if and only if $V_{2} \Perp\left(V_{1}, V_{3}\right)$ or $V_{3} \Perp\left(V_{1}, V_{2}\right)$.

The only difficult part of (3.4) is, in (iii), the implication of $V_{2} \Perp V_{3}$ and $V_{2} \Perp V_{3} \mid V_{1}$. It follows, however, from one of Birch's results (1963, (5.3)) and it completes the proof of Proposition 2.

Remark 1: The hypotheses $V_{2} \Perp V_{3} \mid\left(V_{1}=1\right)$ and $V_{2} \Perp V_{3} \mid V_{1}$, which are of little interest in general $2^{3}$-tables with $V_{1}$ as a response variable, become important in $2^{3}$-tables having marginal independence of treatment $\left(V_{2}\right)$ and site $\left(V_{3}\right)$.

Remark 2: The hypothesis of strongly consistent results in terms of relative risks may be tested with standard methods for odds-ratios (Bishop, Fienberg and Holland, 1975, p. 494).

Remark 3: Equal partial odds-ratios are never collapsible in $2^{3}$-tables with $V_{2} \Perp V_{3}$ and with outcome ( $V_{1}$ ) being marginally dependent on both of treatment $\left(V_{2}\right)$ and site $\left(V_{3}\right)$. This follows from (3.4) (iii) and Proposition 1. The further equivalencies in (3.4) (i) and (ii) imply, together with Proposition 2, that $V_{3}$ has a moderating effect on the odds-ratio in all situations except in the more clear-cut ones, in which either outcome is independent of treatments given sites, (3.4) (i), or outcome is independent of sites given treatments, (3.4) (ii).

Remark 4: If probabilities are estimated for $2^{3}$-tables satisfying $V_{2} \Perp V_{3}$, then tables with $V_{2} \Perp V_{3} \mid\left(V_{1}=i\right)$ for fixed, $i$ are likely to show a reasonable fit to the hypothesis $V_{2} \Perp V_{3} \mid V_{1}$ as well. One can then use the equivalence in (3.4) (iii) to discriminate between the two hypotheses: if statistical tests show $V_{1}$ to depend conditionally or marginally on both of $V_{2}$ and $V_{3}$, then $V_{2} \Perp V_{3} \mid V_{1}$ should be rejected in favour of the less restrictive hypothesis $V_{2} \Perp V_{3} \mid\left(V_{1}=i\right)$ for a fixed $i$.

Proposition 3: In $2^{3}$-contingency tables with outcome ( $V_{1}$ ) depending on sites ( $V_{3}$ ) given treatments $\left(V_{2}\right)$ the hypothesis of no moderating effect of site on the relative risk for $V_{1}=1$ is equivalent to $V_{2} \Perp V_{3}$ and $V_{2} \| V_{3} \mid\left(V_{1}=1\right)$. This result is an immediate consequence of Propositions 1 and 2.

Remark. The combination of the two hypotheses in Proposition 3 does not fit into the log-linear model framework, but it suggests the generalisations treated in the next section.

## 4. THE LACK OF A MODERATING EFFECT ON RELATIVE RISKS

The results of the previous two sections can be exploited to derive nontrivial sufficient conditions for the lack of a moderating effect of one background variable $V_{M}$ on the relative risk in general contingency tables having only nonzero cell probabilities $\Pi$. Let $V_{1}$ be the response, $V_{2}$ a treatment variable, $V_{M}$ be the moderator variable and $V_{3}^{\prime}=V_{3} \cup \ldots \cup V_{M-1}$ the remaining variables, with $I, J, L$ and $K$ categories, respectively. Then the partial relative risks for $V_{1}=1$ under treatment $j\left(V_{2}=j\right)$ compared to treatment $j^{\prime}\left(V_{2}=j^{\prime}\right)$, given $V_{3}^{\prime}$ and $V_{M}$, are

$$
\begin{equation*}
\frac{\Pi_{| | j k l}}{\Pi_{\left[\mid j^{\prime} k l\right.}}=\frac{\Pi_{1 j k l} / \Pi_{+j k l}}{\Pi_{1 j^{\prime} k l} / \Pi_{+j^{\prime} k l}}, \text { all } k, l \tag{4.1}
\end{equation*}
$$

Proposition 4: In an $I \times J \times K \times L$-contingency table with $\Pi_{i j k l}>0$, sufficient conditions for the collapsibility over $V_{M}$ of the partial relative risks (4.1) that are equal for all levels of $V_{M}$ given $V_{3}^{\prime}=k$ are

$$
V_{2} \Perp V_{M} \mid V_{3}^{\prime} \quad \text { or } \quad V_{1} \Perp V_{M} \mid\left(V_{3}^{\prime}, V_{2}\right) .
$$

This follows from the generalisation of the necessary and sufficient condition (A-2) in a $2^{3}$-table to the general case with (4.1) and $j=1, j^{\prime}=2$ :

$$
\Sigma_{l \neq l}\left(\Pi_{1 \mid 2 k l}-\Pi_{1 \mid 2 k l}\right)\left(\Pi_{+i k l} \Pi_{+2 k l^{\prime}}-\Pi_{+2 k l} \Pi_{+1 k l}\right)=0
$$

and the appropriate generalisations of (2.3) (iii) and (3.1) (ii).
Proposition 5: In an $I \times J \times K \times L$-contingency table with $\Pi_{i j k l}>0, V_{M}$ has no moderating effect on the relative risk (4.1)
(i) given $V_{3}^{\prime}=l$ if $\quad V_{2} \Perp V_{M} \mid\left(V_{3}^{\prime}, V_{1}=1\right)$ and $V_{2} \Perp V_{M} \mid V_{3}^{\prime}$,
(ii) irrespective of $V_{3}^{\prime}$ if $V_{2} \Perp\left(V_{M}, V_{3}^{\prime}\right) \mid\left(V_{1}=1\right)$ and $V_{2} \Perp\left(V_{M}, V_{3}^{\prime}\right)$.

For the proof of Proposition 5 (i), note that the two independencies hold if and only if

$$
\Pi_{1 j k l}=\Pi_{1 j k+} \Pi_{1+k l} / \Pi_{1+k+} \quad \text { and } \quad \Pi_{+j k l}=\Pi_{+j k+} \Pi_{++k l} / \Pi_{++k+} .
$$

These imply that, for $j=1, j^{\prime}=2$,

$$
\frac{\Pi_{1 \mid 1 k i}}{\Pi_{1 \mid 2 k i}}=\frac{\Pi_{11 k+} / \Pi_{+1 k+}}{\Pi_{12 k+} / \Pi_{+2 k+}}=\frac{\Pi_{1 \mid k+}}{\Pi_{1 \mid 2 k+}},
$$

so that the partial relative risks (4.1) are equal for all levels $l$ of $V_{M}$ given $V_{3}^{\prime}$ and collapsible over $V_{M}$.

Similarly, note that the two independencies in 5 (ii) are satisfied if and only if

$$
\Pi_{1 j k l}=\Pi_{1 j^{+}} \Pi_{1+k l} / \Pi_{1+++} \quad \text { and } \quad \Pi_{+j k l}=\Pi_{+j++} \Pi_{++k l} .
$$

These give, for $j=1, j^{\prime}=2$,

$$
\frac{\Pi_{1| | k \mid}}{\Pi_{1|2 k|}}=\frac{\Pi_{11++} / \Pi_{+1++}}{\Pi_{12++} / \Pi_{+2++}}=\frac{\Pi_{1 \mid 1++}}{\Pi_{1 \mid 2++}},
$$

so that the partial relative risks (4.1) are equal for all levels of $V_{3}$ and $V_{M}$ and collapsible over $V_{3}^{\prime}$ and $V_{M}$. This completes the proof.

Lemma: In an $I \times J \times K \times L$-contingency table with $\Pi_{i j k t}>0$ following a multinomial distribution for a given total number of observations,
(i) $\quad V_{2} \Perp V_{M} \mid V_{3}^{\prime}$ and $V_{2} \mathbb{L} V_{M} \mid\left(V_{3}^{\prime}, V_{1}\right)$ if and only if in addition either of $V_{1} \Perp V_{2} \mid\left(V_{3}^{\prime}, V_{M}\right)$ or $V_{1} \Perp V_{M} \mid\left(V_{2}, V_{3}^{\prime}\right)$ holds
(ii) $\quad V_{2} \Perp V_{M} \mid V_{3}^{\prime}$ and $V_{2} \Perp V_{M} \mid\left(V_{3}^{\prime}, V_{1}=1\right)$ (i.e. $V_{M}$ has moderating effect on the relative risk (4.1) given the levels of $V_{3}^{\prime}$ ) can be true, if no other independencies are satisfied.

The first part (i) is a special case of Proposition 3 in Wermuth and Lauritzen (1983) and (ii) is proven with the estimated probabilities for the data presented in the next section.

## 5. A SET OF DATA WITH OBSERVED RELATIVE RISKS BEING NEARLY COLLAPSIBLE

From a follow-up investigation by Spielberger on smoking habits of students (Spielberger et al., 1984) observations on 2317 persons are summarised in a $2 \times 3 \times 3 \times 2$-table, where the variables are as follows,
$V_{1}:$ smoking habits of the respondent
$V_{2}$ : older sibling as role model
$V_{3}$ : parents as role model
$V_{4}$ : sex of the respondent.
The first variable is a dichotomous response variable, $V_{2}$ and $V_{3}$ are the main influences of interest and $V_{4}$ is a background variable.

The first two columns in Table 2 show the observed frequencies $n_{1 j k l}$ and $n_{+j k l}=\Sigma_{i} n_{i j k l}$. The third column contains the observed risks to smoke (i.e. for $V_{1}=1$ ), $p_{1 \mid j k l}=n_{1 j k l} / n_{+j k l}$. Those reflect what is supported by standard log-linear analyses, namely that smoking habits of the students $\left(V_{1}\right)$ depend conditionally on all three of $V_{2}, V_{3}, V_{4}$; compare Table 3.
From the hypotheses related to sex $\left(V_{4}\right)$ having no moderating effect on the relative risks to

TABLE 2
Observed and estimated* risks to smoke for 2317 students

| Sex of respondent | Smoking <br> behaviour <br> of parents | Older siblings | Observed frequencies |  | Observed risks to smoke$p_{1 \mid j k l}$ | Estimated risks to smoke$\tilde{p}_{1 \mid j k l}$ | Estimated relative risks |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  |  | $\begin{aligned} & \text { Siblings' } \\ & \text { effects } \end{aligned}$ $\tilde{\tilde{p}}_{1 \mid j k l}$ |  | Parents effects $\tilde{p}_{\left.1 \mid j k^{\prime}\right]}$ |
|  |  |  |  |  |  |  | $P_{1 \mid 2 k i}$ | $\tilde{p}_{1 \mid j 21}$ |
|  |  |  | $n_{i j k l}$ | $n_{+j k t}$ |  |  |  |  |
| female | none smokes | no | 1 | 4 |  | . 250 | . 223 | 1.09 | . 55 |
|  |  | do not smoke | 37 | 172 | . 215 | . 205 | . 58 |  |
|  |  | smoke | 26 | 52 | . 500 | . 545 | 2.66 | . 93 |
|  | one smokes | no | 8 | 18 | . 444 | . 407 | 1.16 |  |
|  |  | do not smoke | 61 | 178 | . 343 | . 351 |  |  |
|  |  | smoke | 103 | 174 | . 592 | . 587 | 1.67 |  |
|  | both smoke | no | 20 | 34 | . 588 | . 466 | 1.00 | 1.14 |
|  |  | do not smoke | 97 | 191 | . 508 | . 465 |  | 1.32 |
|  |  | smoke | 222 | 311 | . 714 | . 750 | 1.61 | 1.28 |
| male | none smokes | no | 1 | 7 | . 143 | . 163 | 1.09 | $\begin{aligned} & .46 \\ & .49 \\ & .77 \end{aligned}$ |
|  |  | do not smoke | 36 | 254 | . 142 | . 149 |  |  |
|  |  | smoke | 21 | 49 | . 429 | . 396 | 2.66 |  |
|  | one smokes | no | 5 | 16 | . 313 | . 354 | 1.16 |  |
|  |  | do not smoke | 48 | 152 | . 316 | . 306 |  |  |  |
|  |  | smoke | 83 | 164 | . 506 | . 511 | 1.67 |  |
|  | both smoke | no | 9 | 32 | . 281 | . 402 | 1.00 | 1.14 |
|  |  | do not smoke | 61 | 169 | . 361 | . 403 |  | 1.32 |
|  |  | smoke | 233 | 340 | . 685 | . 650 | 1.61 | 1.27 |

[^1]TABLE 3
Selected likelihood-ratio chi-squared test results for the data in Table 2

| Variables in the table | Conditional independence given all remaining variables for pair | $\chi^{2}$-value | Degrees of freedom | $p$-value |
| :---: | :---: | :---: | :---: | :---: |
| $\left(V_{1}, V_{2}, V_{3}, V_{4}\right)$ | $\left(V_{1}, V_{2}\right)$ | 150.0 | 12 | 0.000 |
|  | $\left(V_{1}, V_{3}\right)$ | 104.5 | 12 | 0.000 |
|  | $\left(V_{1}, V_{4}\right)$ | 22.9 | 9 | 0.006 |
|  | $\left(V_{2}, V_{3}\right)$ | 222.6 | 16 | 0.000 |
|  | $\left(V_{2}, V_{4}\right)$ | 15.7 | 12 | 0.205 |
|  | $\left(V_{3}, V_{4}\right)$ | 18.8 | 12 | 0.094 |
| $\left(V_{2}, V_{3}, V_{4}\right)$ | $\left(V_{2}, V_{3}\right)$ | 322.6 | 8 | 0.000 |
|  | $\left(V_{2}, V_{4}\right)$ | 7.3 | 6 | 0.291 |
|  | $\left(V_{3}, V_{4}\right)$ | 20.7 | 6 | 0.002 |

smoke (compare Proposition 5), it turns out that only $V_{4} \Perp V_{2} \mid V_{3}$ in the marginal $3 \times 3 \times 2$ table of $\left(V_{2}, V_{3}, V_{4}\right)$ and $V_{4} \Perp V_{2} \mid\left(V_{3}, V_{1}\right)$ in the joint table of $\left(V_{1}, V_{2}, V_{3}, V_{4}\right)$ fit reasonably though not well. These results and the Lemma, however, reveal that both of the hypotheses cannot be satisfied but that instead $V_{4} \Perp V_{2} \mid V_{3}$ and $V_{4} \Perp V_{2} \mid\left(V_{3}, V_{1}=1\right)$ may hold, that is, for known parental smoking patterns ( $V_{3}$ ), sex of the respondent has no moderating effect on the relative risks obtained by comparing situations that differ with respect to the role model older siblings provide.

We proceed by computing smoothed estimates of risks as follows. First, the maximumlikelihood estimates $\hat{m}_{i j k l}$ of the cell frequencies $m_{i j k l}$ are obtained under the independence hypothesis $V_{4} \Perp V_{2} \mid V_{3}$ on the ( $V_{2}, V_{3}, V_{4}$ )-margin (compare Birch, 1963). These are then modified to satisfy the additional hypothesis $V_{4} \Perp V_{2} \mid\left(V_{3}, V_{1}=1\right)$. More precisely we compute for the data at hand,

$$
\begin{align*}
& \hat{m}_{i j k l}=\frac{n_{i j k l} n_{+j k+} n_{++k l}}{n_{+j k l} n_{++k+}}  \tag{i}\\
& \tilde{m}_{1 j k l}=\frac{\hat{m}_{l j k+} \hat{m}_{1+k l}}{\hat{m}_{1+k+}}, \tilde{m}_{2 j k l}=\hat{m}_{+j k l}-\tilde{m}_{1 j k l} \tag{ii}
\end{align*}
$$

(iii) $\tilde{\Pi}_{1 \mid j k l}=\frac{\tilde{m}_{1 j k l}}{\tilde{m}_{+j k l}}$.

The probability estimates $\tilde{\Pi}_{1 ; j k l}$ fit both of the hypotheses $V_{4} \Perp V_{2} \mid\left(V_{3}, V_{1}=1\right)$ and $V_{4} \Perp V_{2} \mid V_{3}$.
The estimates are reasonable in the sense that they provide a good approximation to the data in Table 3 and permit a more condensed description of the effects, since they fit conditions for the lack of a moderating effect of the background variable to the data. The most outstanding result for these data concerns families, in which parents do not smoke ( $V_{3}=1$ ): if a student has older siblings who smoke ( $V_{2}=3$ ), then the risk that he or she ( $V_{4}=1$ or $V_{4}=2$ ) will also smoke is estimated to be $166 \%$ higher than for a student whose older siblings do not smoke ( $V_{2}=2$ ). This is reflected in the relative risk of 2.66 and holds even though the data convey a clear conditional dependence of the smoking behaviour ( $V_{1}$ ) on the sex of the student $\left(V_{4}\right)$ : the risks to smoke are higher for females than for males at all levels of the remaining variables ( $V_{1}, V_{3}$ ). The lack of a moderating effect of $V_{4}$ on the relative risk of $V_{1}=1$ for $V_{2}$ implies that one can estimate the effects of older siblings more stably than others estimated from the four-way table: they are in fact based on observations in a marginal table, that of $V_{1}$, $V_{2}$ and $V_{3}^{\prime}$. However, the statistical properties of our estimates in (4.1) still need to be investigated
and a formal goodness-of-fit test has to be developed. In a forthcoming paper it will be shown how our estimates relate to conditional (Andersen, 1973) and overall maximum-likelihood estimates.

## 6. DISCUSSION

During the last twenty years, the statistical literature on contingency tables strongly emphasised the use of log-linear models for analysing structures. Within this framework marginal independencies are, in general, not easily incorporated (see Birch, 1963). As a consequence, tables of the type discussed in this paper have received little or no attention and their nice properties have not been noticed. Model classes for contingency tables that permit one to impose structure on marginal tables, as well, only seem to have been proposed by Goodman (1973), Wermuth and Lauritzen (1983), Kiiveri, Speed and Carlin (1984) and Lauritzen and Wermuth (1984).

Collapsibility in contingency tables has been defined either for probabilities (Lauritzen, 1982; Asmussen and Edwards, 1983) or for interaction parameters (see Simpson, 1951; Darroch, 1963; Plackett, 1969; Bishop, 1971; Whittemore, 1978). The former proved helpful in answering questions about the equivalence of models. However, the Yule-Simpson paradox can occur when conditions for collapsibility in probabilities are satisfied (examples have been given by Wermuth, 1986). This concept was neither designed to study conditions under which parameters measuring associations, other than the probabilities, may correctly be evaluated in a marginal table, nor is it appropriate for this purpose.

We have shown that conditions for collapsibility of relative risks in tables having a dichotomous response variable involve independencies in the marginal distribution of the variables influencing the response (Proposition 1). The opposite pole to collapsibility of associations or interactions in contingency tables is the Yule-Simpson paradox, and this is caused by strong associations among the influences or regressor variables. The Yule-Simpson paradox can be seen as analogous to problems discussed, in other contexts, under the name of multicollinearity in multiple linear regression (Goldberger, 1964) or as the effects of nonorthogonal factors in analyses of variance (Snedecor and Cochran, 1967). Conditions for the absence of the Yule-Simpson paradox coincide with those under which relative risks possess nice properties: in the case of independent regressors, the marginal relative risk is a simple weighted average of the partial relative risks, equal partial relative risks are always collapsible (3.2), and the hypothesis of equal partial relative risks is equivalent to a conditional independence (3.3).

In view of these results it no longer seem necessary to only approximate relative risks by odds-ratios, as has been proposed by Armitage (1975). One can plan studies in such a way that the resulting data fit hypotheses on marginal independencies perfectly. A decision on this has, of course, to depend on the purpose of the study: if in the target population of the study no marginal independencies are expected, then this can be an important argument against forcing them upon any sample. Many studies are, however, for reasons not discussed here, designed in such a way that a main variable of interest and a background variable behave like independent regressor variables: for instance, all multicentre clinical trials, in which equal numbers of patients are assigned to the treatments within sites, and all multicentre retrospective studies having equal numbers of patients with and without a given disease at all sites. Having equal numbers of patients within sites is just a special case of proportional allocation of observable units to treatments ( $V_{2}$ ) within sites $\left(V_{3}\right)$, and the latter is the empirical counterpart of the marginal independence of $V_{2}$ and $V_{3}$. Given such data, one can test the hypothesis that site has no moderating effect on the relative risk with standard methods (compare Proposition 2).

For more general types of data one can still relate the hypotheses of collapsibility over site and of no moderating effect of site on the relative risk to conditional independencies (Propositions 3, 4 and 5), and these permit us to reconcile seemingly contradictory results in data analysis (Compare the Lemma and Tables 2 and 3).

## ACKNOWLEDGEMENT

I want to thank Peter McCullagh, Paul Holland, Inge and Hanns-Georg Leimer for helpful comments on an earlier version of this paper and a referee, whose concise remarks helped me to strengthen and generalise the results.

## APPENDIX

Here we prove Proposition 1 (ii). We assume again that all probabilities $\Pi_{i j k}$ are positive.
In a $2^{3}$-table with equal partial relative risks for $V_{1}=1$ given $V_{3}$, this relative risk is collapsible over $V_{3}$ if and only if $V_{1} \Perp V_{3} \mid V_{2}$ or $V_{2} \Perp V_{3}$.

Note that the marginal risk for $V_{1}=1$ given $V_{2}=j$ is, for $j=1,2$,

$$
\begin{align*}
\Pi_{1 \mid j+} & =\Pi_{1 j+} / \Pi_{+j+} \\
& =\left(\Pi_{1 \mid j 1} \Pi_{+j 1}+\Pi_{1 \mid j 2} \Pi_{+j 2}\right) / \Pi_{+j+} . \tag{A-1}
\end{align*}
$$

With equal partial relative risks for $V_{1}=1$ given $V_{3}$ we have

$$
\frac{\Pi_{1| | 1}}{\Pi_{1 \mid 21}}=\frac{\Pi_{1 \mid 12}}{\Pi_{1 \mid 22}}
$$

and can express the marginal relative risk, by using ( $\mathrm{A}-1$ ), as

$$
\frac{\Pi_{1 \mid 1+}}{\Pi_{1 \mid 2+}}=\frac{\Pi_{1 \mid 11}}{\Pi_{1 \mid 21}} \frac{\left(\Pi_{1 \mid 21} \Pi_{+11}+\Pi_{1 \mid 22} \Pi_{+12}\right) \Pi_{+2+}}{\left(\Pi_{i \mid 21} \Pi_{+21}+\Pi_{1 \mid 22} \Pi_{+22}\right) \Pi_{+1+}}
$$

This shows that the marginal relative risk equals the partial one if and only if

$$
\left(\Pi_{1 \mid 21} \Pi_{+11}+\Pi_{1 \mid 22} \Pi_{+12}\right)\left(\Pi_{+21}+\Pi_{+22}\right)=\left(\Pi_{1 \mid 21} \Pi_{+21}+\Pi_{1 \mid 22} \Pi_{+22}\right)\left(\Pi_{+11}+\Pi_{+12}\right) .
$$

After rewriting we obtain

$$
\begin{equation*}
\left(\Pi_{1 \mid 21}-\Pi_{1 \mid 22}\right)\left(\Pi_{+11} \Pi_{+22}-\Pi_{+21} \Pi_{+12}\right)=0 \tag{A-2}
\end{equation*}
$$

which is satisfied if and only if

$$
\frac{\Pi_{1 \mid 21}}{\Pi_{1 \mid 22}}=1 \quad \text { or } \quad \frac{\Pi_{+11} \Pi_{+22}}{\Pi_{+21} \Pi_{+12}}=1
$$

These two conditions are, from (2.3) (iii) and (3.1) (ii), seen to be equivalent to $V_{1} \Perp V_{3} \mid\left(V_{2}=2\right)$ and $V_{2} \Perp V_{3}$, respectively. With the assumption of equal partial risks, the first implies $V_{1} \Perp V_{3} \mid V_{2}$ and the proof is complete.

## REFERENCES

Andersen, E. B. (1973) Conditional Inference and Models for Measuring. København: Mentalhygiejnisk Forlag.
Armitage, P. (1975) The use of the cross-product ratio in aetiological survey. In: Perspectives in Probability and Statistics edited by J. M. Gani), pp. 343-355. London, New York: Academic Press.
Asmussen, S. and Edwards, D. (1983) Collapsibility and response variables in contingency tables. Biometrika, 70, 567-578.
Birch, M. W. (1963) Maximum likelihood in three-way contingency tables. J. R. Statist. Soc. B, 25, 220-233.
Bishop, Y. M. M. (1971) Effects of collapsing multidimensional contingency tables. Biometrics, 27, 545-562.
Bishop, Y. M. M., Fienberg, S. E. and Holland, P. W. (1975) Discrete Multivariate Analysis: Theory and Practice. Cambridge, Mass. and London: MIT Press.
Currie, I. and Korabinski, A. (1984) Some comments on bivariate regression. The Statistician, 33, 283-293.
Darroch, J. N. (1962) Interactions in multifactor contingency tables. J. Royal Statist. Soc. B, 24, 251-263.
Dempster, A. P. (1971) An overview of multivariate data analysis. J.Multivar. Anal., 1, 316-46.
Ekholm, A. (1985) A recursion formula for the log linear parameters of a collapsed contingency table. Research Report 53, Department of Statistics, University of Helsinki.
Golberger, A. S. (1964) Econometric Theory. New York: Wiley.

Goodman, L. A. (1973) The analysis of contingency tables, when some variables are posterior to others: a modified path analysis approach. Biometrika, 60, 179-192.
Hill, A. B. (1962) Statistical Methods in Clinical and Preventive Medicine. New York: Oxford Univ. Press.
Kiiveri, H., Speed, T. P. and Carlin, J. B. (1984) Recursive causal models, J. Australian Math. Soc. A, 36, 30-52.
Lauritzen, S. L. (1982) Lectures on Contingency Tables, 2nd printing. Aalborg: Aalborg Univ, Press.
Lauritzen, S. L. and Wermuth, N. (1984) Mixed interaction models. Research report, Aalborg Universitets Center (ISSN-0106-0791).
Plackett, R. L. (1969) Multidimensional contingency tables: a survey of models and methods. Bull. Int. Statist. Inst., 43, 133-141.
Simpson, E. H. (1951) The interpretation of interaction in contingency tables. J. R. Statist. Soc. B, 13, 238-241.
Snedecor, G. W. and Cochran, W. G. (1967) Statistical Methods, 6th edition. Ames: Iowa State University Press.
Spielberger, C. D., Jacobs, G. A., Crane, R. S. and Russell, S. F. (1983) On the relation between family smoking habits and the smoking behaviour of college students. Int. Rev. Appl. Psychol., 32, 54-69.
Wermuth, N. (1976) Analogies between multiplicative models in contingency tables and covariance selection. Biometrics, 32, 95-108.
(1980) Linear recursive equations, covariance selection, and path analysis. J. Amer. Statist. Ass., 75, 963-97.
(1986) Implications of the Yule-Simpson paradox for simple multicentre studies. In: Research Report 86-1, Stochastik und verwandte Gebiete, University of Mainz. (ISSN 0177-0098).
Wermuth, N. and Lauritzen, S. L. (1983) Graphical and recursive models for contingency tables, Biometrika, 70, 537-552.
Whittemore, A. S. (1978) Collapsibility of multidimensional contingency tables. J. R. Statist. Soc. B, 40, 328-340.
Yule, G. U. (1900) On the association of attributes in statistics. Phil. Trans. Ser A., 194, 257-319.
Zedeck, S. (1971) Problems with the use of "moderator" variables. Psychol. Bull., 76, 295-310.


[^0]:    $\dagger$ Address for correspondence: Professor N. Wermuth, Psychologisches Institut, Johannes Guttenberg-Universität, Postfach 3980, D-6500 Mainz, Federal Republic of Germany.

[^1]:    * Estimates as defined in (4.1)

