

Residual-based Shadings for Visualizing (Conditional) Independence

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Visualizing independence by association and mosaic plots.

Enhancements: diverging color palette based on HCL color space combined with visualization of the result of a significance test.

Extensions to visualizing conditional independence.

Key Words: Association plots; Conditional inference; Contingency tables; HCL colors; HSV colors; Mosaic plots; Permutation tests.

1. INTRODUCTION

independence in 2-way tables

association and mosaic displays

colors and color spaces

somewhere: mention Augsburg stuff, visualization of log-linear models via mosaics ([Theus and Lauer 1999](#); [Hofmann 2003, 2001](#)).

Association plots as residual plots for log-linear models ([Meyer, Zeileis, and Hornik 2003](#)), especially in coplot or trellis layout.

2. INDEPENDENCE IN 2-WAY TABLES

In this section, the basic tools for testing and visualizing independence in 2-way tables are briefly reviewed. For illustration, a data set about treatment and improvement of patients with rheumatoid arthritis from [Koch and Edwards \(1988\)](#) is used. The data set is also discussed in [Friendly \(2000\)](#) and the subset of the 59 female patients from the study is given in Table 1.

2.1 TESTS

To fix notations, we consider a 2-way contingency table with cell frequencies $\{n_{ij}\}$ for $i = 1, \dots, I$ and $j = 1, \dots, J$ and row and column sums $n_{i+} = \sum_j n_{ij}$ and $n_{+j} = \sum_i n_{ij}$ respectively. For convenience the number of observations is denoted $n = n_{++}$. Given an underlying distribution with theoretical cell probabilities π_{ij} , the null hypothesis of independence of the two categorical variables can be formulated as

$$H_0 : \pi_{ij} = \pi_{i+}\pi_{+j}. \quad (1)$$

The expected cell frequencies in this model are $\hat{n}_{ij} = n_{i+}n_{+j}/n$. The probably best known and most used measure of discrepancy between observed and expected values are the Pearson residuals

$$r_{ij} = \frac{n_{ij} - \hat{n}_{ij}}{\sqrt{\hat{n}_{ij}}}. \quad (2)$$

The most convenient way to aggregate the $I \times J$ residuals to one test statistic is their squared sum

$$X^2 = \sum_{i,j} r_{ij}^2, \quad (3)$$

Table 1: Treatment and improvement among 59 patients with rheumatoid arthritis.

		Improved		
		None	Some	Marked
Treatment	Placebo	19	7	6
	Treated	6	5	16

because this is known to have an unconditional limiting χ^2 distribution with $(I-1)(J-1)$ degrees of freedom under the null hypothesis. This is the well-known χ^2 test which is typically introduced at the very beginning of the chapter about independence in 2-way tables in statistics textbooks (see e.g., [Agresti 2002](#)).

But the sum of squares is not the only plausible way of capturing deviations from 0 in the Pearson residuals. There are many other conceivable functionals $\lambda(\cdot)$ which lead to reasonable test statistics $\lambda(\{r_{ij}\})$: one which is particularly suitable for identifying the cells which cause the dependence (if any) is the maximum of the absolute values

$$M = \max_{i,j} |r_{ij}|. \quad (4)$$

Given a critical value c_α for this test statistic, all residuals whose absolute value exceeds c_α violate the hypothesis of independence at significance level α ([Mazanec and Strasser 2000](#), Ch. 7). Thus, the interesting cells causing the dependence can easily be identified.

Furthermore, an important reason for using the unconditional limiting distribution for the X^2 statistic (3) was the closed form result for the distribution. Recently, with the improving performance of computers, conditional inference (or permutation tests)—carried out either by simulation or by computation of the (asymptotic) permutation distribution—have been receiving increasing attention ([Pesarin 2001](#); [Strasser and Weber 1999](#)). For testing the independence hypothesis (1), using a permutation test is particularly intuitive due to the permutation invariance (given row and column sums) of this problem. Consequently, all results in this paper are based on conditional inference performed by simulating from the permutation distribution of test statistics of type $\lambda(\{r_{ij}\})$.

Other measures of discrepancy (e.g., deviance residuals) could, of course, also be used instead of the the Pearson residuals $\{r_{ij}\}$. But as the ideas discussed here extend straightforwardly to that situation, we do not discuss it in detail.

For the arthritis data from Table 1, both tests indicate a clearly significant dependence of improvement on treatment: the sum-of-squares statistic from Equation (3) is $X^2 = 11.296$ with a p value of $p = 0.0032$, and the maximum statistic from Equation (4) is $M = 1.87$ with a p value of $p = 0.0096$. Both p values have been computed by drawing a sample of size 5000 from the permutation distribution under independence.

2.2 VISUALIZATIONS

Two well-established visualization techniques for independence in 2-way tables are mosaic plots and association plots. Both are suitable to bring out departures of an observed table $\{n_{ij}\}$ from the expected table $\{\hat{n}_{ij}\}$ in a graphical way. The latter focuses on the visualization of the Pearson residuals r_{ij} (under independence) while the latter primarily displays the observed frequencies n_{ij} .

Mosaic plots ([Hartigan and Kleiner 1981](#)) can be seen as an extension of grouped bar charts where width and height of the bars show the relative frequencies of the two variables: a mosaic plot simply consists of a collection of tiles whose sizes are proportional to the observed cell frequencies as shown in the left panel of Figure 1. A rectangle corresponding to 100 percent of the observations is first split horizontally with respect to the treatment frequencies and then vertically with respect to the conditional improvement frequencies. This shows that there have been more placebo than treated patients with no improvement and vice versa for marked improvement. This strategy of splitting with respect to conditional frequencies given all previous variables can also directly be

used for visualizing multi-way tables (see [Hofmann 2003](#), for an overview of how to construct and read mosaic displays).

Association plots ([Cohen 1980](#)) visualize the table of Pearson residuals: each cell is represented by a rectangle that has (signed) height proportional to the corresponding Pearson residual r_{ij} and width proportional to the square root of the expected counts $\sqrt{\hat{n}_{ij}}$. Thus, the area is proportional to the raw residuals $n_{ij} - \hat{n}_{ij}$. The association plot for the arthritis data is shown in the right panel of Figure 1 which leads to the same interpretation as the mosaic plot: there are more placebo patients with no improvement and fewer with marked improvement than expected under independence—vice versa for the treated patients.

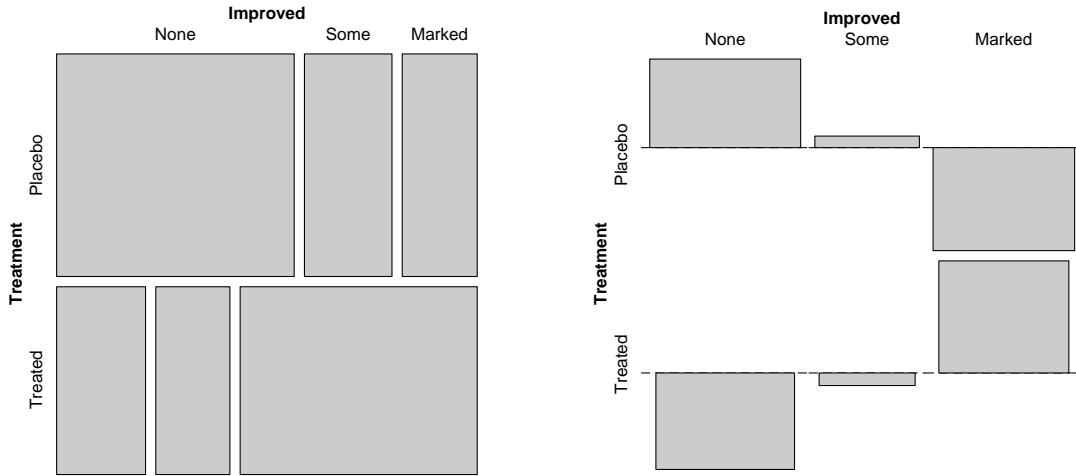


Figure 1: Classic mosaic and association plot for the Arthritis data.

3. RESIDUAL-BASED SHADINGS

remarks: Friendly 1994

A first extension by [Friendly \(1994\)](#) uses a color coding of the tiles to visualize deviations (residuals) from a given log-linear model fitted to the table, that is, from the expected frequencies under independence. This approach does not only work in 2-way tables but also in log-linear models fitted to multi-way tables. In this extension, positive and negative signs of the residuals are coded by rectangles with solid and dashed borders respectively. Furthermore, residuals exceeding an absolute value of 2 are shaded light blue and red respectively, those that even exceed an absolute value of 4 are shaded with full saturation. The heuristic behind this shading is that the Pearson residuals are approximately standard normal which implies that the highlighted cells are those with residuals *individually* significant at approximately the 5% and 0.01% level. But the main purpose of the shading is not to visualize significance but the *pattern* of deviation from independence ([Friendly 2000](#), p. 109). In particular, this shading does not provide a visualization of the maximum test (4) as in general it is unknown to which significance level α the values $c_\alpha = 2$ and 4 correspond for the table being visualized.

3.1 COLORS

currently: chosen based on HSV or HLS color spaces, hue (red and blue) codes sign of residuals, and saturation the absolute size in three steps, value currently unused (always set to full value)

disadvantages: - perceptually not uniform because the three perceptual dimensions (hue, lightness, saturation) are not properly mapped to the three dimensions of the color space and hence are confounded ([Brewer 1999](#)) in particular, saturation is not uniform across different hues.

- can lead to color-caused optical illusions in statistical graphs ([Cleveland and McGill 1983](#))
- flashy fully saturated HSV colors are good for drawing attention to a plot, but hard to look at for a longer time, hence mosaic displays using these colors are harder to interpret
- white is not very suitable as a neutral color, grey conveys neutrality or uninterestingness much better

Alternatives: - [Munsell \(1905\)](#) introduced color notation for balanced colors - Commission Internationale de l'Éclairage (CIE) introduced two perceptually based color spaces CIELAB and CIELUV where the latter is preferred for emissive color technologies such as computer displays.

- **ColorBrewer.org** ([Harwer and Brewer 2003](#)) is an online tool for selecting different sets of colors, based on similar ideas. It includes qualitative, sequential and diverging palettes. - [Ihaka \(2003\)](#) discusses how CIELUV colors can be used for choosing colors for statistical graphics such as barplots. He recommends to use polar coordinates in the CIELUV space—called HCL space, using hue, chroma and luminance—and suggests a strategy for choosing qualitative schemes based on these.

We discuss how similar ideas can be used for deriving a diverging HCL scheme that is suitable for visualizing residuals in association and mosaic displays.

HCL - irregular double cone, i.e., chroma/luminance plane looks rather different for different hues - choose two hues that look similar in the chroma/luminance plane to code the sign of the residuals. Also use red and blue at full chroma for large residuals and use a light gray (instead of a white at the top of the luminance scale) as the neutral color for small residuals, interpolate between. - for non-significant results, clearly reduce the amount of color, i.e. the maximum chroma used.

Advantages - based on a perceptually uniform color space - device independent

3.2 SIGNIFICANCE

The extensions of [Friendly \(1994\)](#) to mosaic plots provide substantial improvement of the original mosaic plots and enhanced them from a plot for contingency tables to a visualization technique for log-linear models (and therefore also for certain independence problems). However, it has two major drawbacks: First, the significance level for the hard-coded critical values 2 and 4 is usually unknown. Second, the colors for (light) blue and red used in the Michael Friendly's SAS implementation and also in the base implementation in R are not device-independent and do not provide homogeneous saturations over different colors and copier proofness. We suggest some ways to overcome these drawbacks.

As pointed out in Section 2.1, the critical values for the maximum statistic M from (4) can be derived from the permutation distribution. Instead of the hard-coded values 2 and 4, the particular critical values, e.g., at the levels $\alpha = 0.1$ and 0.01 , for the table to be visualized could be used. Hence, exactly those residuals causing the (potential) dependency within the table are highlighted. At the moment, these critical values are derived in `vcd` by simulation of the underlying distribution.

The implementation of Friendly mosaic plots in SAS uses (by default) colors in the HLS (Hue–Luminance–Saturation) color space with hues for blue and red, full saturation, and varying luminance for lighter colors. The current R implementation is based on colors in the HSV (Hue–Saturation–Value) space where decreasing the saturation from 1 towards 0 generates very similar colors as increasing the luminance for HLS colors from 0.5 towards 1. As an alternative to those HLS and HSV colors, the device-independent HCL (Hue–Chroma–Luminance) color space ([Ihaka 2003](#)) could be used which also provides homogeneous saturations over different colors and copier proofness. The resulting association plot (using critical values for $\alpha = 0.1$ and 0.01) for the Bundesliga data can be seen in Figure ???. It indicates very clearly that there is no evidence for rejecting

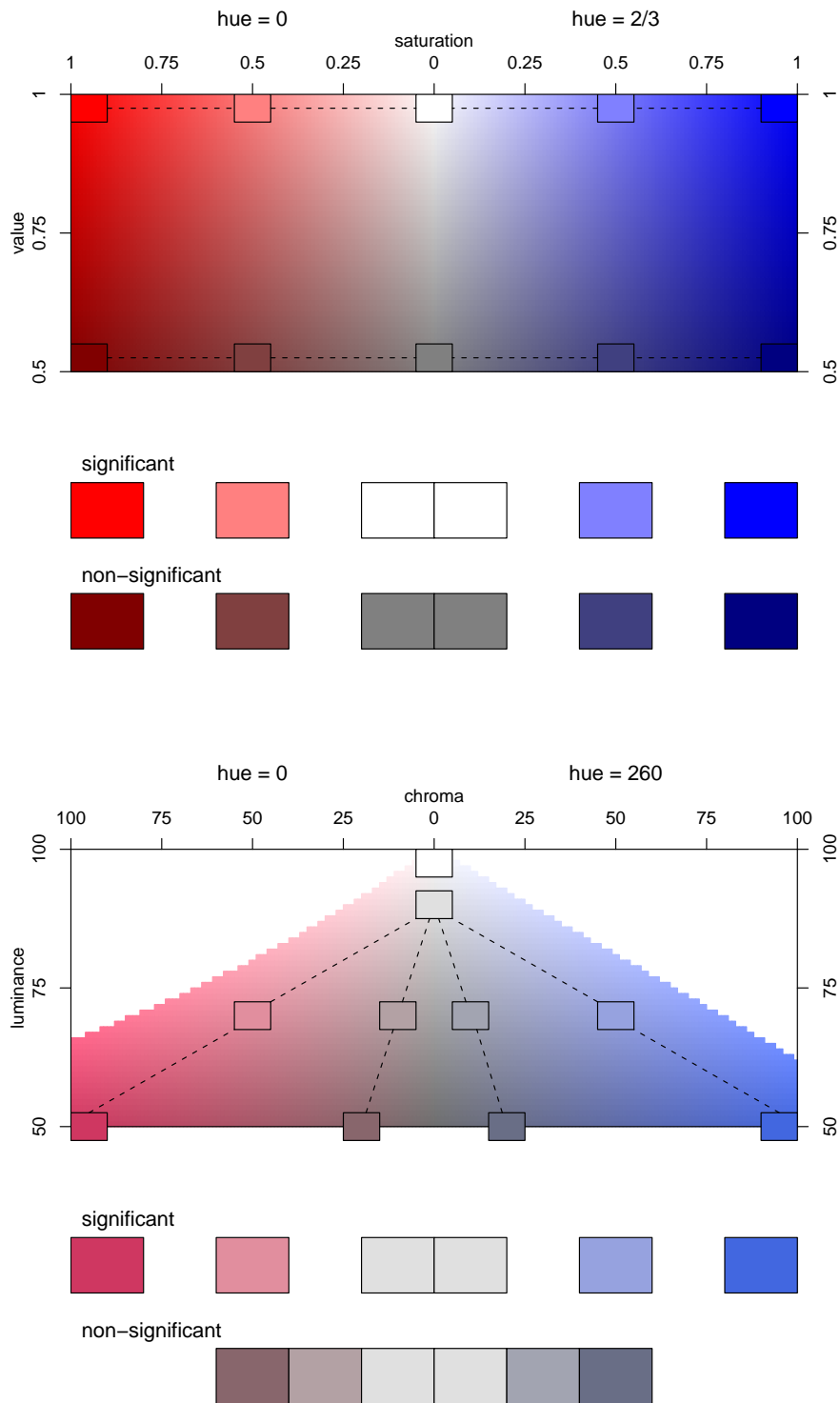


Figure 2: Extended shading in HSV and HCL space.

the hypothesis of independence in this table. Thus, there seems to be neither an advantage for the home team nor for the away team.

To illustrate what a ‘postive’ example looks like we give the association plot for the well-known Hair-Eye-Color data set (available in base R) which cross-tabulates the hair and eye color and gender of 592 statistics students: Figure ?? shows the association plot for the 328 female students. It is easy to see that the hypothesis of independence is rejected at 1% level by the maximum test (4) due to higher frequencies of students with hair and eye color black/brown and blond/blue and lower frequencies of blond/brown and brown/blue than would be plausible under independence. Furthermore, it can also be seen that the residual corresponding to the black/blue cell is significant at the 10% level.

We have seen that this approach performs very well when the test statistic which should be visualized is the maximum statistic (4). However, it is difficult to extend this approach to general tests for independence, in particular in the HCL space, where the ranges of chroma and luminance are not independent. In contrast the HSV space has the advantage that all three dimensions hue, saturation and value have the range $[0, 1]$ and can be varied independently. Therefore, a different approach than the one described above could be to use the hue for the sign of the residuals as before, the saturation for the absolute size of the residuals, and the value as an indicator for the significance of some test statistic (only using the full value when the overall test for independence rejects the null hypothesis). The resulting mosaic plots under this paradigm are depicted in Figure ??. Again, it can clearly be seen that despite some large residuals there is no evidence against independence for the Bundesliga data, but that the null hypothesis has to be rejected for the Hair-Eye-Color data.

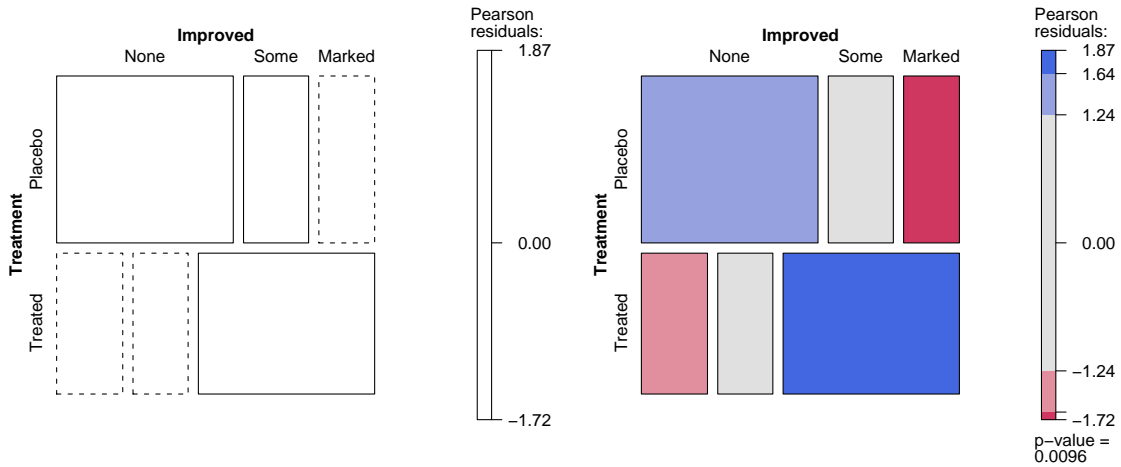


Figure 3: Mosaic plot for the Arthritis data with Friendly shading (left) and extended shading (right).

4. EXTENSIONS TO CONDITIONAL INDEPENDENCE

UCB admissions data (Bickel, Hammel, and O’Connell 1975) double max: 3.134, p-value = 0.0004 max Chisq: 17.248, p-value = 0.0006 sum Chisq: 19.938, p-value = 0.0034 simulated from 5000 repetitions

Friendly (1999) discusses grouping similar to coplots (conditioning plots) (Cleveland 1993) that lead to trellis graphics (Becker, R.A., Cleveland, W.S. and Shyu, M. “The Visual Design and Control of Trellis Display”, Journal of Computational and Graphical Statistics).

Independence problems do not only occur in 2-way tables, although that is an important special case, but they are also important in tables of higher dimensionality and can follow much more complex patterns. These are again defined based on the underlying table of theoretical cell probabilities ($\pi_{ijk\dots}$) with more than two dimensions. Models of interest include the null hypotheses of conditional independence:

$$\pi_{ijk\dots} = \pi_{i|k\dots}\pi_{j|k\dots}$$

Classical non-graphical methods for these problems include the χ^2 test, Fisher’s exact test, the Cochran-Mantel-Haenzel test (for $2 \times 2 \times K$ -tables), and the analysis of log-linear models for more complex settings.

As an example, two natural ways to use the visualization techniques described in the previous sections would be to use (Trellis-like) conditioning plots or pairs plots (like mosaic matrices) to visualize these more complex patterns of independence.

Two ideas for the problem of conditional independence are briefly outlined here and illustrated using the famous admissions data of the University of California at Berkley (UCB) which is available in base R. In this data, the question whether there is gender discrimination at the UCB leads to the result that although women seem to be disadvantaged at the aggregated level there is no gender discrimination conditioned on the department—with the very exception of one department in which women are *more* likely to be admitted than would be plausible under independence. Exactly this is illustrated in the conditioning association plot in Figure ??.

Similarly, the same data can be visualized using a mosaic matrix where a conditional independence model is fitted in each plot (see Figure ??).

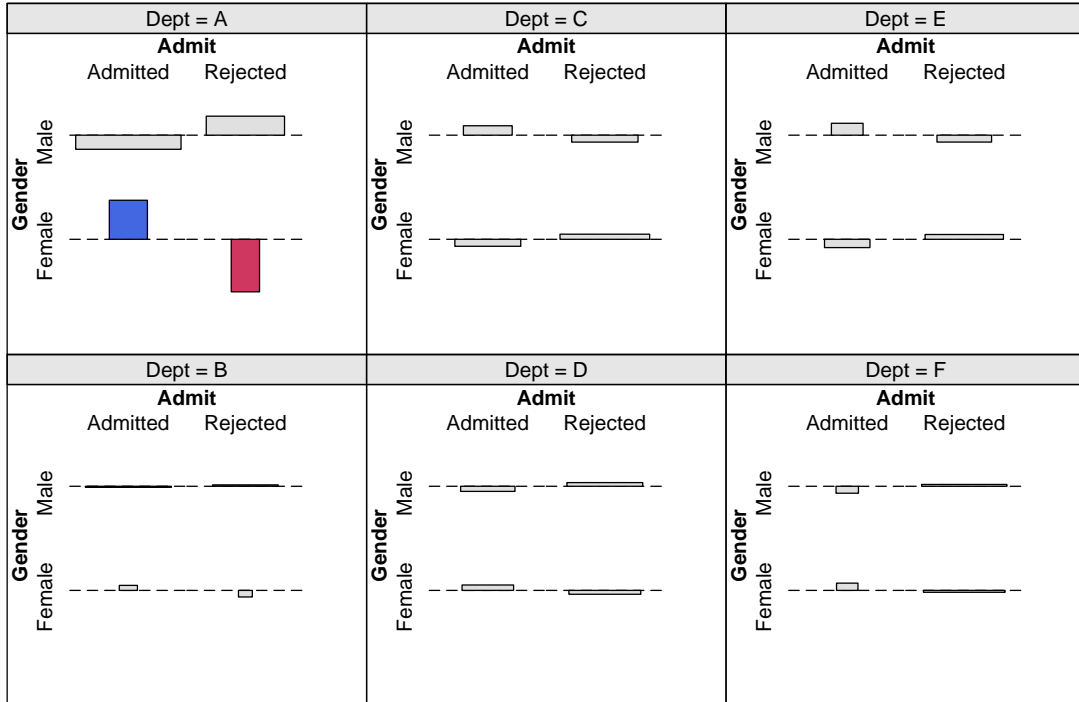


Figure 4: Conditional association plot for UCB admissions.

5. CONCLUSIONS

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