This manuscript is now published as:

de Cheveigné, A (2021). *Harmonic Cancellation - A Fundamental of Auditory*Scene Analysis. **Trends in Hearing**, 25, https://doi.org/10.1177/23312165211041422.

Please refer to and cite that version.

# **Harmonic Cancellation - a Fundamental of Auditory Scene Analysis**

Alain de Cheveigné(1, 2, 3)

#### **AUTHOR AFFILIATIONS:**

- (1) Laboratoire des Systèmes Perceptifs, UMR 8248, CNRS, France.
- (2) Département d'Etudes Cognitives, Ecole Normale Supérieure, PSL, France.
- (3) UCL Ear Institute, United Kingdom.

#### **CORRESPONDING AUTHOR:**

Alain de Cheveigné, Audition, DEC, ENS, 29 rue d'Ulm, 75230, Paris, France.

# Acknowledgements

This work was supported by grants ANR-10-LABX-0087 IEC, ANR-10-IDEX-0001-02 PSL, and ANR-17-EURE-0017. Mickael Deroche, John Culling and Josh McDermott made useful comments on a previous version, and Maria Chait and Israel Nelken offered useful advice. The manuscript also benefitted greatly from comments of two anonymous reviewers and the Editor, Chris Plack.

**Abstract** 

This paper reviews the hypothesis of harmonic cancellation according to which

an interfering sound is suppressed or canceled on the basis of its harmonicity (or

periodicity in the time domain) for the purpose of Auditory Scene Analysis. It

defines the concept, discusses theoretical arguments in its favor, and reviews ex-

perimental results that support it, or not. If correct, the hypothesis may draw on

time-domain processing of temporally-accurate neural representations within the

brainstem, as required also by the classic Equalization-Cancellation (EC) model

of binaural unmasking. The hypothesis predicts that a target sound corrupted by

interference will be easier to hear if the interference is harmonic than inharmonic,

all else being equal. This prediction is borne out in a number of behavioral studies,

but not all. The paper reviews those results, with the aim to understand the incon-

sistencies and come up with a reliable conclusion for, or against, the hypothesis

of harmonic cancellation within the auditory system.

**keywords:** Pitch perception, auditory scene analysis, segregation, harmonicity,

harmonic cancellation

3

# Introduction

Our environment is cluttered with sound sources, but to act effectively we must focus on one or a few and ignore the others. This is hard because the mixing process, by which sounds from the various sources add up before entering the ears, cannot be undone. We usually do not know the mixing matrix (i.e. the delays and gains applied to each source before adding) and, even if we did, that matrix is generally not invertible. Recovering individual sources is thus *impossible* except in very simple cases. Nonetheless, we sometimes feel that we can follow an individual source, for example a voice within a conversation, or an instrument within an ensemble, as if it were alone. The ability to make sense of a complex acoustic scene in terms of individual sources is known as Auditory Scene Analysis (Bregman, 1990).

Auditory Scene Analysis is sometimes discussed as a process of "grouping" elements (e.g. partials) to form sources or objects (Bregman, 1990), for example according to Gestalt principles. However, such "elements" are conceptual rather than operational. While sinusoids and clicks serve well as synthesis parameters, it may not be possible to extract them from the sound due to theoretical limits (e.g. time-frequency uncertainty tradeoff, Gábor, 1947) and physiological limits (e.g. temporal and frequency resolution of cochlear analysis, Moore & Glasberg, 1983; Plack & Moore, 1990). If they cannot be accessed, postulating that they can be grouped is perhaps misleading.

Fortunately, perfect isolation of each source is usually not necessary. According to the principle of *unconscious inference* (Helmholtz, 1867; Kersten, Mamassian, & Yuille, 2004), we need only to recover enough information to infer the presence or nature of a target. Regularities within the world, internalized as models within the perceptual system, allow us to fill in missing parts. This process, which manipulates incomplete information "under the hood", provides us with the

illusion of perceiving each object just as if true unmixing had taken place. Information about the source is partial but, thanks to inference, it appears to us that it is complete (al Haytham, 1030; Hatfield, 2002; Imbert, 2020).

For this to work, it is essential that the sensory representation be stripped of the influence of background objects. If not, a different background might lead to a different percept, defeating the goal of perceiving the target as if it were in isolation. In other words, the sensory representation should be made *invariant* to the presence of interfering sources. This is analogous to invariance with respect to intra-class variability in Pattern Classification (Duda, Hart, & Stork, 2012).

Several aspects of auditory processing might contribute to this goal. If target and background differ by their spectral content, *cochlear filtering* can be used to split sensory input into channels dominated by the target, distinct from those that reflect the background. Discarding the latter then yields a representation that is invariant to the presence of the background – albeit incomplete because of the missing channels. Likewise, if target and background occur at different points in time, *temporal resolution* properties of the auditory system (Moore, Glasberg, Plack, & Biswas, 1988; Plack & Moore, 1990) can be used to discard time intervals contaminated by the background.

Putting both elements together, the target can be "glimpsed" within spectro-temporal gaps of the background (Cooke, 2006). The glimpsed "pixels" of the time-frequency representation are handed over to subsequent processing together with a mask to indicate their position. Discarded pixels are not merely set to zero: they are given *zero weight* (Cooke, Morris, & Green, 1997). Spectro-temporal glimpsing has been proposed in speech processing applications (D. Wang, 2008; D.-L. Wang & Brown, 2006), and to account for human perceptual abilities and derive predictive measures of intelligibility (e.g. Best, Roverud, Baltzell, Rennies, & Lavandier, 2019; Josupeit, Schoenmaker, Par, & Hohmann, 2020).

Binaural disparity is another potentially useful cue. In addition to head shadow effects that produce favorable target-to-masker ratios within certain frequency channels at either ear (Grange & Culling, 2016), perception benefits from binaural interaction, which is commonly understood to follow the well-known Equalization Cancellation (EC) model (Durlach, 1963), and its extensions (e.g. Akeroyd, 2004; Breebaart, van de Par, & Kohlrausch, 2001; Culling & Summerfield, 1994). Signals at each ear are differentially time-shifted and scaled ("equalization"), and then subtracted one from the other ("cancellation") to suppress interaurally-coherent sound from a competing source. The internal time shift and scale factor are tuned to match the interfering source. The EC model is assumed to involve temporally accurate neural patterns processed by specialized neural circuitry within the auditory brainstem (Joris & van der Heijden, 2019; Tollin & Yin, 2005).

To summarize this viewpoint, Auditory Scene Analysis entails cancelling and/or ignoring irrelevant features of the sensory input, and matching the remainder to an internal model to produce a reliable percept. The process draws on spectro-temporal analysis within the cochlea, complemented by neural time-domain signal processing within the brain, to provide the brain with a rich – albeit incomplete – representation within which a target can be "glimpsed". The glimpses are then interpreted according to a Helmholtzian inference process.

The remainder of this paper asks whether this process can be extended to include, as a cue, the harmonic (periodic) structure of interference such as a competing talker. So-called "double-vowel" experiments found that vowels mixed in pairs are easier to identify if their fundamental frequencies ( $F_0$ s) differ (Assmann & Summerfield, 1994; Brokx & Nooteboom, 1982; Culling & Darwin, 1993; McKeown, 1992), suggesting that harmonic structure somehow assists segregation. Furthermore, it appears that this effect is driven mainly by the harmonicity of the *background*, e.g. the competing vowel (de Cheveigné, McAdams, & Marin,

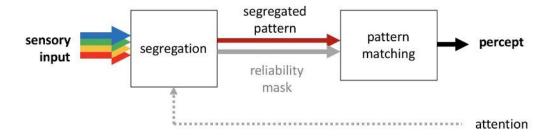


Figure 1: Segregation and matching. Sensory input is stripped of correlates of interfering sources, and the selected pattern, possibly incomplete, is passed on for pattern-matching (or model-fitting), together with a mask that indicates which parts are missing or unreliable. Initial stages are under attentional control.

1997; Lea, 1992; Summerfield & Culling, 1992). This is the harmonic cancellation hypothesis.

To set the stage, I assume a "segregation module" that works hand in hand with a "pattern-matching" module (Fig. 1). The segregated sensory pattern (red arrow) is accompanied by a "reliability mask" (gray arrow) to assist matching of a pattern that is incomplete or distorted by the segregation process. Sensory representations might consist of a spectral profile (e.g. place-rate representation), or a temporal, or place-time pattern. Examples of the latter are a matrix of autocorrelation functions (ACF), one per channel (autocorrelogram), or the sum over channels of these ACFs (summary autocorrelation function, SACF) (Licklider, 1959; Lyon, 1984; Meddis & Hewitt, 1992). The flow of sensory information in this schema is purely bottom-up: the only top-down influence is attentional control (dotted arrow). Top-down transfer of a sensory-like pattern is also conceivable ("schema-driven" segregation), but not considered here.

We want to know whether harmonic cancellation is instantiated in the auditory system, but it is often easier to reason in terms of the acoustic waveform, for clarity and to distinguish theoretical from implementation limits: if a principle fails in abstract terms, consideration of biological constraints is premature. That said,

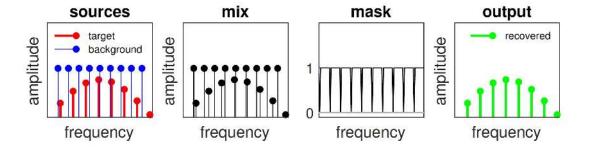


Figure 2: Harmonic cancellation in the idealized frequency domain. Left: line spectra of a "target" sound (red) and a "background" (blue). Next to left: mixture. Next to right: harmonic mask with zeros at all harmonics of background. Right: recovered target.

references to "cochlear filtering" or "neural processing" will sometimes creep into the discussion without warning. I beg your patience when this occurs.

# **Harmonic Cancellation - Possible Mechanisms**

How might harmonic cancellation be implemented? This section investigates several hypotheses, including frequency-domain, time-domain, and hybrid models. A later section will ask which – if any – is used by the auditory system. The busy reader might want to read about *Frequency Domain* and *Time Domain* models, then skip to the Psychophysics section and come back for details as needed. There are also interesting things to be found in the Appendix.

#### Frequency Domain

Conceptually, harmonic cancellation is straightforward: just zero all spectral components at multiples of  $F_0=1/T$  where T is the period of the background, as in Fig. 2 (Parsons, 1976; Stubbs & Summerfield, 1988). Target components emerge intact (right panel), except in the event, vanishingly unlikely in this ideal-

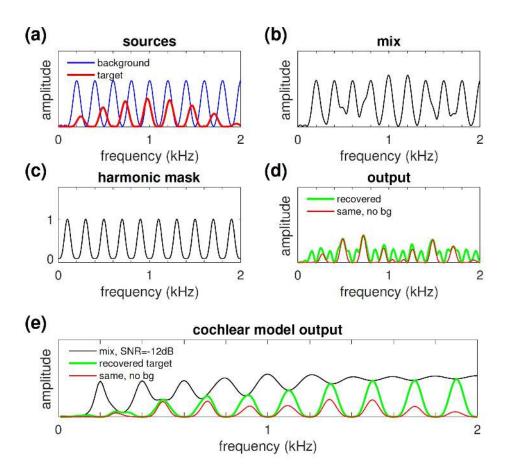


Figure 3: Harmonic cancellation in the frequency domain using a short-term Fourier representation, or a filter bank. (a) 238 Hz target (red) and 200 Hz background (blue) analyzed by a filter bank with 100 Hz resolution, (b) mixture, (c) harmonic mask, (d) target recovered from mixture (green), and same in the absence of the background (thin red), (e) same analysis but using a filter bank with non-uniform frequency resolution. Filter bandwidth depends on center frequency (CF) according to estimates of cochlear frequency resolution from Moore and Glasberg (1983) as implemented by Slaney (1993).

ized world, that a target component falls on the harmonic series of the background.

A practical implementation, however, needs to deal with two issues: one is limited frequency resolution of the spectral representation, the other is the spectral widening expected when analyzing a time-limited and/or non-stationary signal. Figure 3 (a) shows short-term amplitude spectra of two harmonic sounds, a 200 Hz "background" with a flat spectral envelope (blue), and a weaker 238 Hz "target" with a broad peak centered at 1 kHz (red).

This spectral transform has limited frequency resolution (or, equivalently, infinite resolution but the signals are time-limited, in this case eight cycles of a 200 Hz fundamental, shaped with a Hanning window). When target and masker are mixed, here with a target-to-masker ratio (TMR) of -12 dB, the spectrum of the mix (Fig. 3 b, black) is almost entirely dominated by the background (Fig. 3 a, blue). This differs radically from the idealized picture of Fig. 2.

If we multiply the spectrum with a harmonic mask with zeros at the harmonics of the background (Fig. 3, c), we obtain a "recovered" spectral pattern (d, green) very different from the true target (a, red). Two terms contribute to this difference. One is multiplicative distortion from the masking procedure (compare d, red to a, red), the other is additive distortion due to the incompletely-cancelled background (compare d, green to d, red). The former can, in principle, be taken into account by a pattern-matching stage if it has access to the nature of that distortion, for example via the gray arrow in Fig. 1. The latter is more serious because it is unknown and cannot be compensated for, and because it implies that we miss our goal of invariance with respect to the background. The shape of the harmonic mask (Fig. 3 c) affects the balance between error terms but a different mask would not yield a radically different result. The contrast between Fig. 2 (conceptual model) and Fig. 3 (implementation) is sobering.

Spectral resolution is critical. Cochlear filters are narrower, on a linear fre-

quency scale, at low than at high center frequencies (CF) (Fig. 3 e). From this figure it would seem that low-frequency target features might be recovered, but perhaps not high-frequency (compare green and thin red). This illustration used a bank of gammatone filters (Slaney, 1993) with equivalent rectangular bandwidths (ERBs) from psychophysical estimates (Moore & Glasberg, 1983). If cochlear filters were narrower (e.g. Shera, Guinan, & Oxenham, 2002; Sumner et al., 2018) a wider frequency range might be recoverable (not shown), but resolution would still be limited if the stimulus were short or non-stationary.

In summary, frequency-domain cancellation requires (a) a spectral representation with resolution sufficient to cancel background partials while retaining enough of the target to support pattern matching, (b) an estimate of the background period T, and (c) a pattern-matching process that tolerates distortion of target spectral patterns. How to estimate the background period is discussed in the Appendix (*Period Estimation*).

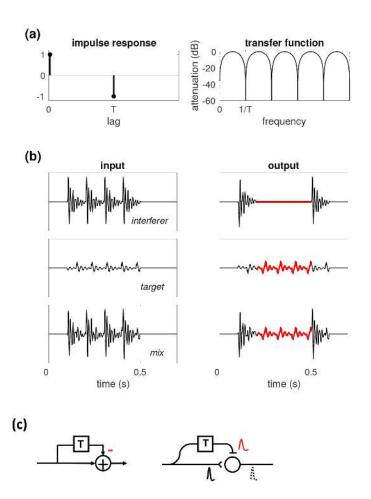


Figure 4: Harmonic cancellation in the time domain. (a) Impulse response of the cancellation filter (left) and corresponding magnitude transfer function (right). (b) Input (left) and output (right) of the cancellation filter for the background 100 Hz vowel/a/(top), target 132 Hz vowel/e/(middle), and mixture at TMR=-12 dB (bottom). (c) Schematic of a circuit implementing the cancellation filter (Eq. 1) (left) and neural circuit with similar function (right). A spike on the direct pathway (black) is transmitted unless it coincides with a spike on the delayed pathway (red). The delay can be applied to the positive/excitatory input, instead of negative/inhibitory, with equivalent results.

#### Time Domain

Harmonic cancellation can also be implemented in the time domain by a sim-

ple filter with impulse response

$$h(t) = \delta_0(t) - \delta_T(t) \tag{1}$$

where T is the period of the interfering sound and  $\delta_T$  is the Kronecker delta function translated to T (Fig. 4 a, left). The filtered version of a signal s(t) is simply s(t) - s(t - T). The magnitude transfer function of this filter has deep dips at all harmonics of 1/T (Fig. 4 a, right).

Figure 4 (b) shows a background vowel stimulus /a/ with fundamental 100 Hz (top), a weaker target vowel /i/ with fundamental 132 Hz (middle), and their mixture (bottom), before (left) and after (right) filtering with a cancellation filter with lag T equal to the period of the background vowel. The response consists of initial and final one-period glitches, separated by a short steady-state portion, in red. The steady-state portion is zero for the background (top). For the target, it is a distorted version of the target waveform (compare middle right, red, to middle left). For the mixture, it is the same as for the target alone (compare middle right, red, to bottom right, red). In other words, this part of the pattern is *invariant with respect to the presence of a background* of period T, which is what we need. This contrasts with frequency-domain cancellation for which none of the recovered pattern was background-invariant.

In summary, time-domain cancellation requires (a) a time-domain signal representation such that Eq. 1 can be implemented, (b) an estimate of the background period T (see Appendix,  $Period\ Estimation$ ), (c) a pattern matching process capable of selecting the intervals of perfect cancellation, and compensating for distortion of the target within these intervals.

#### Hybrid Models

A hybrid model combines spectral and temporal processing, for example cochlear

filter bank analysis followed by time-domain harmonic cancellation within the brainstem. There is a rich literature based on this idea for the purpose of auditory modeling and sound processing applications (e.g. Assmann & Summerfield, 1990; Lyon, 1983, 1988; Meddis & Hewitt, 1992; Weintraub, 1985). A benefit of the filter bank is that TMR varies across channels, some favoring the target and others the background (Fig. 5 a), which may be useful if the dynamic range of temporal processing is limited.

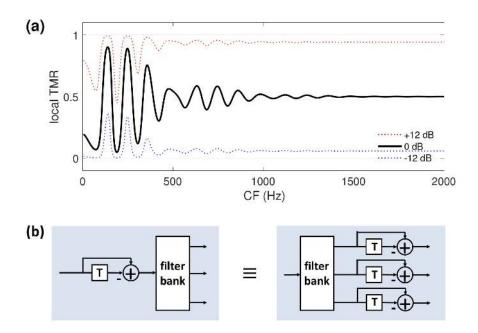


Figure 5: (a) TMR within each channel of a model cochlear filter bank for an input consisting of a 124 Hz harmonic target mixed with a 100 Hz harmonic background with overall TMR=0 dB (black), -12 dB (dotted blue), or +12 dB (dotted red). Thanks to the filter bank, the TMR is enhanced in certain channels within which the target can be "glimpsed". (b) Linear operations can be swapped. Filtering the signal before the filter bank is equivalent to applying the same filter to each channel after the filter bank.

It is worth remembering that linear, time-invariant operators can be swapped: a time-domain cancellation filter applied to the acoustic waveform can instead be applied to each channel after filtering: the result is the same (Fig. 5 b). Cochlear

filtering and transduction are both non-linear and non-stationary (e.g. adaptation), but the "equivalence" of Fig. 5 (b) may nonetheless be useful conceptually. I review briefly here a selection of hybrid schemes for harmonic cancellation, described in detail in the Appendix (*Hybrid Models*). In brief:

- Hybrid Model 1: Cancellation-enhanced spectral patterns. A timedomain cancellation filter is applied to each channel of the cochlear filter bank, resulting in sharper selectivity and cleaner spectral patterns for pattern matching.
- Hybrid Model 2: Channel rejection on the basis of periodicity. Channels dominated by the background periodicity are discarded, and the remaining channels are used to form a time-domain pattern for pattern matching, as in the concurrent vowel identification model of Meddis and Hewitt (1992).
- Hybrid Model 3: Cancellation filtering of selected channels. As in Hybrid Model 2, channels dominated by the background are discarded, and channels dominated by the target are left intact. In contrast to Hybrid Model 2, channels with intermediate TMR are processed by a cancellation filter. The result is used for time-domain pattern matching.
- Hybrid Model 4: Channel-specific cancellation filter. The parameter T
   of the cancellation filter can differ between channels, in contrast to other
   models that use the same T for all channels. The result is used for time domain pattern matching.
- Hybrid Model 5: Synthetic delays. The "synthetic delay" mechanism of de Cheveigné and Pressnitzer (2006) is used to implement the relatively long delays T required by the temporal model of harmonic cancellation. The result is used for time-domain pattern matching.

• Hybrid Model 6: Logan's theorem. This is not a specific model but a processing principle. A narrowband signal can be reconstructed perfectly from its zero crossings (and hence also from its half-wave rectified version) (Logan, 1977). This implies that, despite the non-linearities, the temporal model can be implemented after transduction as if it were applied to the acoustic waveform (the theorem does not say how).

These examples illustrate how peripheral filtering and temporal processing might work hand-in-hand to enhance a spectral model (Hybrid model 1) or a temporal model (Hybrid Models 2-6) of harmonic cancellation. To summarize, a wide variety of mechanisms can implement harmonic cancellation: spectral, temporal and hybrid.

#### Alternatives to Harmonic Cancellation

It is important to consider alternatives: to the extent that they are viable, the case for harmonic cancellation is weaker. Other aspects of the spectral structure of the target or background might support segregation, even in situations that seem to implicate harmonic cancellation.

#### Harmonic Enhancement.

According to this hypothesis, the harmonic structure of a *target* sound allows its extraction from a background. The idea is attractive: it fits with the Auditory Scene Analysis credo that components of a sound must be "grouped" together, here on the basis of harmonicity, to form a coherent "object" that can be distinguished from other parts of the scene (Bregman, 1990). It is satisfying to hypothesize that voiced speech might be "engineered" for this purpose through evolution (e.g. Popham, Boebinger, Ellis, Kawahara, & McDermott, 2018).

The mechanisms just reviewed can be re-purposed for enhancement. For example, the mask in Fig. 2 can be made to *select* target harmonics rather than reject

background harmonics. Likewise, replacing the minus by a plus in Eq. 1, and setting T to the period of the target, yields a *harmonic enhancement filter*:

$$h(t) = \delta_0(t) + \delta_T(t). \tag{2}$$

Enhancement and cancellation seem symmetric one of the other, but they have rather different properties. Enhancement requires the period of the target, but this is hard to estimate when TMR is small, which is unfortunately when segregation is most necessary. Cancellation works well in that situation. An enhancement filter provides only a limited boost in TMR (6 dB for the simple filter of Eq. 2) in contrast to cancellation that can reject the masker perfectly, at least in principle. A larger boost would require a longer impulse response (as explained in Appendix A of de Cheveigné, 1993, courtesy of Jean Laroche), but this might not be practical for a non-stationary signal such as speech. Anticipating, behavioral results also don't favor the enhancement hypothesis.

Incidentally, the term "harmonic enhancement" appears in other contexts with a different meaning: perceptual enhancement of one harmonic of a complex when it is turned on or off (e.g. Hartmann & Goupell, 2006). Hopefully no confusion will result from this overloading of the terminology.

Spectral Glimpsing. Between the lines of a harmonic spectrum are gaps where target components might be glimpsed (Deroche, Culling, & Chatterjee, 2013; Guest & Oxenham, 2019), and this might conceivably account for the benefit observed when a background is harmonic rather than inharmonic. Figure 5 (a) shows how individual channels in the low frequency region can preferentially reflect one source or the other, as long as partials are not too close. The spectral-glimpsing hypothesis glosses over the question of how target channels are distinguished from background channels. In that, it differs from Hybrid Model 2 above.

#### Waveform Interactions.

The sinusoidal waveforms of two or more partials can interact within a channel of a filter bank to produce a complex "beat" pattern. This can occur between partials of the same sound (with a rate equal to the fundamental if the sound is harmonic) or partials of different sounds. The patterns that result are quite diverse (static summation, slow fluctuations, rapid beats, etc.), and they depend in a complex way on several parameters (frequencies, levels, filter shapes). The "waveform interactions" hypothesis is thus ill-defined unless further specified.

From slow to fast: phase-dependent summation of same-frequency partials constitutes a potential confound in experiments that include a "zero  $\Delta F_0$ " condition (de Cheveigné, 1999c). Slow beats between closely-spaced partials from different sounds cause the short-term spectrum to cycle between shapes that might favor perception of one or the other sound, either because it momentarily resembles that of one of the sounds in isolation, or because temporal contrast effects enhance important spectral features (Assmann & Summerfield, 1994; Culling & Darwin, 1994; Summerfield, Foster, Gray, & Haggard, 1981). Faster beats might evoke a sensation of roughness signaling the presence of a target (Treurniet & Boucher, 2001), or the spectral location of such beats might provide cues to its spectral features (e.g. the location of a formant peak, or the boundary between formants of different sounds). Conversely, the *lack* of beats at a rate slower than  $F_0$  (or the perceptual correlate of this lack, "smoothness") could signal the absence of a target, or the spectral location of channels dominated by harmonics of a single sound. Finally, the absence of any modulation at  $F_0$  implies that the channel is dominated by a single partial, as in the phenomenon of "synchrony capture" which might signal the position of a formant peak of a successfully isolated sound (Carney, Li, & McDonough, 2015; Maxwell, Richards, & Carney, 2020).

Interaction of more than two harmonics produces a phase-dependent beat pat-

tern that is more deeply sculpted for certain phase relations, such as cosine, or "Klatt" phase that approximates natural phonation with a glottal pulse within each period. Valleys between pulses might then allow a target to be glimpsed for a favorable alignment, as might occur if sounds of different  $F_0$  are mixed (the pitch period asynchrony hypothesis, PPA, Summerfield & Assmann, 1991).

Beat patterns might be exploited to group channels by correlation (Fishman & Steinschneider, 2010; Hall, Haggard, & Fernandes, 1984; Shamma, Elhilali, & Micheyl, 2011; Sinex, Henderson Sabes, & Li, 2002; Sinex & Li, 2007) or, alternatively, beat rates in the  $F_0$  range might be compared across channels (Roberts & Bregman, 1991; Roberts & Brunstrom, 2003; Treurniet & Boucher, 2001). This requires the existence of some mechanism to analyze beat patterns and quantify their rates (see *Modulation Filter Bank* below).

Beat amplitude depends non-monotonically on the amplitude of sources within the stimulus, and the shape of the beat pattern is phase-dependent (for three or more partials). Beat rate affects perceptual salience (e.g. roughness) non-monotonically, and the rate itself may depend non-monotonically on  $F_0$  difference, depending on which partials happen to be close. Finally, each channel has its own pattern of beats. For these reasons, a "waveform interaction hypothesis" is hard to delineate and test (which does not imply that it is incorrect).

#### Modulation Filter Bank.

An influential idea is that cochlear filtering and transduction are followed by analysis by a *modulation filter bank* within the auditory system (Dau, Kollmeier, & Kohlrausch, 1997; Jepsen, Ewert, & Dau, 2008; Joris, Schreiner, & Rees, 2004; Kay & Matthews, 1972; Stein, Ewert, & Wiegrebe, 2005; Viemeister, 1979). Conceptually, this seems rather like reproducing internally an operation (spectral analysis) that is already carried out in the cochlea. A major difference, however, is that it occurs after *demodulation* of each output of the peripheral filter bank (non-

linearity followed by smoothing), which makes it primarily sensitive to features of the waveform envelope, and less sensitive to carrier phase. The concept makes most sense when applied to slow fluctuations (e.g. below  $\sim 30$  Hz), but models have been proposed with channels up to  $\sim 500$  Hz, capitalizing on the smooth transition between neural coding of fine structure at low frequencies and of envelope at higher frequencies (Joris et al., 2004). A modulation filter bank applied to each peripheral channel results in a center frequency  $\times$  best modulation frequency pattern that can be collapsed across channels to obtain a "summary modulation spectrum". One could imagine a frequency-domain harmonic cancellation model applied to this "internal spectrum". However, most estimates of modulation filter width are rather wide (quality factor  $Q \approx 1$ ), which makes this idea unlikely to work given the issues mentioned earlier.

Alternatively, the 2-D pattern could be used to tag channels for the purpose of segregation (Ewert & Dau, 2000; Meyer, Plante, & Berthommier, 1997). One might consider implementing this modulation filter bank using cancellation filters, which would result in a model similar to the hybrid models reviewed previously, a major difference being the demodulation step which renders the model sensitive to envelope periodicity rather than (or in addition to) waveform periodicity.

#### In Summary.

Multiple models have been put forward to explain how the harmonic structure of sounds within an acoustic scene can be used to analyze the scene and attend to particular sources. Some fit the definition of harmonic cancellation, others do not. The next section reviews psychophysical evidence in favor – or against – this hypothesis and its alternatives.

# **Psychophysics**

Detection Benefits from  $\Delta F_0$ .

When presented with a mixture of two vowels, subjects more often report that they hear two vowels if the  $F_0$ s differ (Arehart, Rossi-Katz, & Swensson-Prutsman, 2005; Arehart, Souza, Muralimanohar, & Miller, 2011; de Cheveigné, McAdams, & Marin, 1997; McPherson, Dolan, et al., 2020). Likewise, when presented with a harmonic tone with one partial mistuned, they may detect the partial as "standing out" as a separate sound (Moore, Glasberg, & Peters, 1986; Moore, Peters, & Glasberg, 1985). Such a mistuned target tone can be detected at  $\sim -15$  dB relative to a harmonic masker, whereas against a noise background the threshold is  $\sim 15$  dB higher (Micheyl, Bernstein, & Oxenham, 2006). In each of these examples, background harmonicity seems to affect how many sources are heard. An interpretation, in the context of harmonic cancellation, is that a single entity is perceived if cancellation is perfect, and multiple entities if it leaves a residual.

#### *Discrimination and Identification Benefit from* $\Delta F_0$ .

Mistuning one partial of a harmonic complex allows it to be matched to a pure tone (Hartmann, McAdams, & Smith, 1990), implying not only that this "second sound" is detectable, but also that its frequency can be accessed. Subjects are more likely to identify both vowels of a concurrent pair if their  $F_0$ s differ (Arehart et al., 2011; Assmann & Summerfield, 1994; Brokx & Nooteboom, 1982; Chalikia & Bregman, 1993; Culling & Darwin, 1993; McKeown, 1992; Scheffers, 1983; Shackleton, Meddis, & Hewitt, 1994; Summerfield & Assmann, 1991; Zwicker, 1984). The pattern of results is similar across studies: poor performance (albeit well above chance) for  $\Delta F_0$ =0, rapid improvement up to about one semitone, fol-

lowed by a plateau and possibly a dip at the octave. To create the  $\Delta F_0$ =0 condition with continuous speech, the voices must be re-synthesized on a monotone, or one voice given the same  $F_0$  track as the other, so that  $\Delta F_0$ s remain the same throughout the presentation. With that manipulation, a similar benefit of non-zero  $\Delta F_0$  is obtained (Brokx & Nooteboom, 1982; Leclère, Lavandier, & Deroche, 2017).

Improved performance with  $\Delta F_0 \neq 0$  is taken to reflect a harmonicity-based segregation mechanism that fails when  $F_0$ s are the same, and indeed, identification is less good if both voices are whispered (Lea, 1992), or inharmonic (de Cheveigné, McAdams, & Marin, 1997). This brings up the question as to whether each voice benefits from its harmonic structure, that of its competitor, or both. To answer that question, voices must be parametrized individually, and responses tallied separately. It cannot be answered if the performance metric is "both correct" (Brokx & Nooteboom, 1982; Scheffers, 1983; Summerfield & Assmann, 1991), or if both voices are made inharmonic at the same time (Popham et al., 2018).

#### Background Harmonicity is Important.

In "double vowel experiments", listeners give two answers on each trial, but it has been noted that one constituent (the "dominant" vowel) is usually identified regardless of  $\Delta F_0$ , whereas identification of the other depends on  $\Delta F_0$  (McKeown, 1992; McKeown & Patterson, 1995; Zwicker, 1984). "Dominance" is phoneme- and subject-dependent, but this can be overridden by changing the relative level of the vowels, in which case the  $\Delta F_0$  effects are mainly observed for the weaker (smaller amplitude) vowel (Arehart et al., 2005; de Cheveigné, Kawahara, Tsuzaki, & Aikawa, 1997; McKeown, 1992). This is congruent with the harmonic cancellation hypothesis, in that estimation of the harmonic structure of the background should be easy when the target is weak. However, it could also simply result from a reduced ceiling effect for the more challenging, weaker vowel.

With the  $\Delta F_0 \neq 0$  condition as a starting point, performance decreases if the

competing vowel is whispered (Lea, 1992) or made inharmonic (de Cheveigné, McAdams, & Marin, 1997), regardless of whether the target is harmonic or not. This too is consistent with the harmonic cancellation hypothesis. Similar results are reported for connected speech: Steinmetzger and Rosen (2015) found that speech reception thresholds were up to 11 dB lower for periodic than aperiodic maskers, while Deroche, Culling, Chatterjee, and Limb (2014b) reported a 4 dB elevation in speech reception threshold (SRT) for inharmonic vs harmonic maskers. Incorporating harmonic cancellation within a predictive model of speech intelligibility improved its fit to experimental data (Prud'homme, Lavandier, & Best, 2020).

Gockel, Moore, and Patterson (2002) found that the threshold for detecting noise in a harmonic masker was 11 to 14 dB lower than the converse, and Gockel, Moore, and Patterson (2003) found a similar result for loudness. This suggests that a harmonic masker might be less potent than a noise masker, as expected from harmonic cancellation. As mentioned earlier, Micheyl et al. (2006) found that a harmonic complex tone (HCT) was easier to detect within a background consisting of another HCT than within noise, and Klinge, Beutelmann, and Klump (2011) found a lower threshold for detection of a tone embedded in (but mistuned from) a harmonic rather than inharmonic or noise background (see also Oh & Lutfi, 2000).

All these results are consistent with harmonic cancellation. However, harmonic cancellation is not exclusive of other mechanisms, and one might expect the auditory system to use several or all if they are effective. The next section reviews evidence for harmonic enhancement.

#### Target Harmonicity is Less Important

The idea that harmonicity ensures that a sound does not "fall apart into a sea of individual harmonics" is seducing (Popham et al., 2018), but studies that tried

to demonstrate an advantage of target harmonicity for segregation have met with mixed results. As noted earlier, in double-vowel experiments the benefit of a  $\Delta F_0$  is greatest for weak targets, and measurable for TMR as low as -25 dB (Arehart et al., 2005; de Cheveigné, Kawahara, et al., 1997; McKeown, 1992). Estimating the  $F_0$  of a target that weak would be challenging. Replacing a voiced target by a whispered target does not impair intelligibility, regardless of whether the competitor is voiced or whispered (Lea, 1992), nor does randomly perturbing its harmonics to make it inharmonic (de Cheveigné, McAdams, & Marin, 1997). Modulating the  $F_0$  of target speech in the presence of reverberation disrupts its periodicity, but Culling, Summerfield, and Marshall (1994) found no effect on SRTs (see also Deroche & Culling, 2011b).

For continuous speech, it has been hypothesized that target harmonicity (one aspect of "temporal fine structure", TFS) could aid glimpsing within a spectro-temporally modulated noise, by tagging time-frequency regions that are voiced. However, a direct test of this hypothesis gave negative results (Shen & Pearson, 2019). There is however some evidence that continuity of target  $F_0$  helps to connect information over time, or reduce informational masking if target and masker  $F_0$  ranges are non-overlapping (Darwin & Bethell-Fox, 1977).

A difficulty in testing the enhancement hypothesis is that manipulation of the target might affect its intelligibility independently of any segregation effect. Whispered speech is reportedly less intelligible than voiced speech (Ruggles, Freyman, & Oxenham, 2014), and reverberation, which disrupts harmonicity of an intonated target, also degrades intelligibility (Deroche & Culling, 2011b). Manipulating  $F_0$  (monotonizing, transposing, or inverting the  $F_0$  track) may also affect intrinsic intelligibility (Binns & Culling, 2007; Deroche, Culling, Chatterjee, & Limb, 2014a; Guest & Oxenham, 2019). Such effects might conceivably offset the benefits of harmonic enhancement, making them unmeasurable, so the best we can

say is that we lack strong evidence in favor of that hypothesis.

#### An Intriguing Exception: Target Pitch

In contrast to results just reviewed, a target within a noise background *is* easier to detect if it is harmonic than inharmonic (McPherson, Grace, & McDermott, 2020). This inconsistency is resolved if we reflect that a harmonic target is likely detected in noise on the basis of its pitch (Gockel, Moore, Plack, & Carlyon, 2006; Hafter & Saberi, 2001; Scheffers, 1984), which is probably more salient if the sound is harmonic. If frequency discrimination in noise relies on a pitch percept, it too should benefit from target harmonicity, as found by McPherson, Grace, and McDermott (2020). Thus, we cannot with confidence attribute such benefits to enhanced *segregation* as opposed to an enhanced *pitch percept*.

It is also intriguing that the pitch of a target is easier to discriminate if mixed with a noise background rather than a harmonic background (Micheyl et al., 2006), opposite to what we expect of harmonic cancellation (indeed, the same sounds were easier to *detect* within a harmonic background than a noise background). It would seem that background harmonicity interferes with target pitch, possibly in a way similar to the phenomenon of pitch discrimination interference (PDI) (Gockel, Carlyon, & Plack, 2009; Micheyl, Keebler, & Oxenham, 2010). That interference is not absolute: the pitch of a mistuned partial may be heard within a harmonic background (Hartmann & Doty, 1996; Hartmann et al., 1990), and individual tones may be heard within a chord (Graves & Oxenham, 2019), consistent with skills found in competent musicians.

#### Is the Benefit Explained by Spectral Glimpsing?

Several results seem consistent with this hypothesis. The benefit of  $\Delta F_0$  to vowel identification is mainly limited to the region of resolved partials (Culling & Darwin, 1993), and it improves with a higher background  $F_0$  at which partials are

more widely spaced (Deroche et al., 2013; Deroche, Culling, Chatterjee, & Limb, 2014a). Guest and Oxenham (2019) found that removing the even harmonics of a masker reduced masking of a target placed one octave above, also consistent with glimpsing within the large gaps between background partials of odd rank.

However, Deroche et al. (2013); Deroche, Culling, Chatterjee, and Limb (2014a, 2014b) argued that the larger gaps that arise when a masker is made inharmonic should *reduce* masking, contrary to their results. A possible explanation is that cancellation and glimpsing are both involved (Deroche, Culling, Chatterjee, & Limb, 2014b), consistent with Hybrid models 2 or 3.

## Is the Benefit Explained by Waveform Interactions?

As pointed out earlier, waveform interaction comes in multiple forms, and it is not always clear which version of the hypothesis is implied when it is invoked. One difficulty, common to many versions, is that the non-monotonic dependency of beat amplitude on component amplitudes implies that the magnitude (and spectral locus) of beat-dependent cues should show non-monotonic variations with level, whereas identification usually varies monotonically with TMR. Another challenge is that  $F_0$ -based segregation seems to benefit mostly partials of low rank, for which, thanks to resolvability, the distribution over channels of high-amplitude beats is likely sparse (Deroche, Culling, & Chatterjee, 2014).

Phase effects attributable to PPA were found at 50 Hz, but not at 100 Hz or higher (de Cheveigné, McAdams, & Marin, 1997; Deroche et al., 2013; Deroche, Culling, & Chatterjee, 2014; T. Green & Rosen, 2013; Summerfield & Assmann, 1991, but see Summers and Leek 1998). Furthermore, reverberation should scramble the phase relations required by PPA, whereas it does not affect segregation unless  $F_0$  is modulated (Culling, Hodder, & Toh, 2003; Culling et al., 1994; Deroche & Culling, 2011b).

Culling and Darwin (1994) attributed effects of small  $\Delta F_0$  to the ability to

shop for favorable spectral patterns among those offered by slow beats. Random starting phase should reduce this benefit due to the haphazard temporal alignment of beat patterns, but, de Cheveigné, McAdams, and Marin (1997) found that the  $\Delta F_0$  benefit did not depend on the phase pattern (random vs sine) of either target or background. The slow-beat hypothesis was further tested by de Cheveigné (1999c), again with limited support. The reader should refer to those two papers for a detailed discussion of several forms of the waveform interactions hypothesis. Given the diversity, it is hard to rule out that some form of waveform interaction contributes to segregation. Indeed, harmonic cancellation itself could be construed as a mechanism to exploit a particular form of waveform interaction specific to harmonically-related partials.

#### The Special Case of Maskers with Frequency-shifted or Odd-order Harmonics

In experiments that require detecting (or matching the pitch of) a mistuned partial of rank n within a harmonic complex of fundamental  $F_0$ , the subject likely attends to channels with a center frequency close to  $nF_0$ . The task might then be hampered by the presence, within those channels, of neighboring harmonics, in particular harmonics of rank n-1 and n+1. A cancellation filter tuned to  $F_0$  would suppress those unwanted harmonics, but it would also suppress the target *unless it is mistuned*. We would thus expect performance to improve with mistuning, as indeed is observed (Hartmann et al., 1990; Moore et al., 1986).

However, Roberts and Brunstrom (1998) found a similar result when the background series had been made inharmonic by shifting all partials by the same amount  $\Delta f$ , in which case partials are regularly spaced but harmonicity is disrupted. This suggests that *spectral regularity*, rather than harmonicity, might be the driving factor, which would put in doubt the harmonic cancellation account. However, that proposal hinges on the existence of a mechanism to detect spectral regularity: Roberts and Brunstrom (2001) doubted the existence of a dictionary of

shifted-harmonic templates.

An alternative is that harmonic cancellation is applied *locally* within peripheral channels, for example based on Hybrid model 4 (analogous to what has been proposed for the binaural EC model, Akeroyd, 2004; Breebaart et al., 2001; Culling & Summerfield, 1994). Specifically: the shifted partials  $(n-1)F_0 + \Delta f$  and  $(n+1)F_0 + \Delta f$  can be approximated with harmonics of rank n-1 and n+1 of a harmonic series of fundamental  $F_0(1+\Delta f/n)$ . A cancellation filter tuned to that series would approximately cancel the closest offending background partials (more distant ones are attenuated by cochlear filtering). The nth zero of that filter falls at  $nF_0 + \Delta f$ , i.e. it fits the "spectral regularity" template invoked by Roberts and Brunstrom (1998), which would explain why they found that "mistuning" a partial from that position makes it easier to detect or match. An array of such CF-dependent cancellation filters, each tuned to an "equivalent  $F_0$ " equal to  $F_0(1+\Delta f/f_c)$  would attenuate a shifted-harmonic complex across all channels, allowing "mistuning" relative to that spectrally regular (but inharmonic) pattern to be detected.

This reasoning can be extended to the case of a background harmonic complex with only odd harmonics of  $F_0$ , as it is equivalent to a series of harmonics of  $2F_0$  each shifted by  $\Delta f = -F_0$ . This series can be cancelled *perfectly* by a cancellation filter tuned to  $F_0$ , or *approximately*, within each peripheral channel, by a cancellation filter tuned near  $2F_0$  as just described. The reason for considering the latter is that it requires a shorter delay, which is relevant if there is a penalty on longer delays as has been suggested in the context of pitch perception (Bernstein & Oxenham, 2008; de Cheveigné & Pressnitzer, 2006; Moore, 2003). An array of cancellation filters, each tuned to  $2F_0(1 + F_0/f_c)$ , would spare anything that does not fit the series of odd harmonics, in particular an even-numbered harmonic. If so, it might explain why a single even-numbered harmonic embed-

ded among odd-numbered harmonics is "heard out" more easily than any of the odd-numbered partials (Roberts & Bregman, 1991), and similar explanation might underlie the benefit for identification of a speech target of removing even harmonics of the masker (Guest & Oxenham, 2019) mentioned earlier. This question is revisited in the Discussion.

#### *In Summary*

A body of evidence agrees with the hypothesis that harmonic cancellation assists auditory scene analysis, complementing the well-known benefits of peripheral frequency analysis. Dissenting results are sparse. The alternative hypothesis of harmonic enhancement, while attractive, garners little experimental support. Harmonic cancellation raises a number of issues that are discussed further in the Appendix. These include *period estimation* (necessary to apply cancellation), the relations between *correlation and cancellation*, analogies with the well-known *EC model* of Durlach, *pattern matching* with missing data, potential *anatomical and physiological substrates*, and the possible synergy between *cochlear filtering* and *neural filtering*.

## **Discussion**

Periodicity (or harmonicity) – and its perceptual correlate, pitch – have long captured the attention and imagination of thinkers and scientists (Micheyl & Oxenham, 2010). A periodic sound within the right parameter range evokes a salient percept that is long-lasting in memory (McPherson, Dolan, et al., 2020), is robust to masking by noise (Hafter & Saberi, 2001; McPherson, Grace, & McDermott, 2020), and supports fine discrimination (e.g. Micheyl & Oxenham, 2010). However, the idea that a sound "falls apart" unless it is harmonic does not withstand a bit of reflection. A one-period tone pulse seems unitary without the aid of har-

monicity, meaningless at that duration. A harmonic tone of longer duration may sound unitary, but so does noise which lacks harmonicity. An alternative proposition is that the percept evoked by a sound is unitary by default, and that "multiplicity" is inferred from the accumulation of evidence in favor of additional sources. A complex with a mistuned harmonic initially sounds like a single object but, given time and encouragement, a subject might detect something amiss and interpret it as an additional source. The process requires time (Hartmann et al., 1990; McKeown & Patterson, 1995; Moore et al., 1985), and is harder if the background is made inharmonic (Roberts & Brunstrom, 2003; Roberts & Holmes, 2006). Thus, one could argue, the harmonic nature of one part of the stimulus makes it easier to detect the presence of other parts. From this perspective, harmonicity of a source may contribute to a percept of *multiplicity* for mixtures in which it participates, rather than to its own unity.

That background harmonicity is crucial comes as a surprise, as it suggests that segregation must rely on an adventitious quality of the environment. Also surprising is that target harmonicity has only a minor role, as it goes against the attractive idea that communication sounds are "engineered" through evolution to be harmonic for resilience. It does make sense, however, when one realizes that cancellation works well (and enhancement poorly) at low TMR, which is when segregation is most needed. Infinite TMR improvement can be achieved, in principle, for very short stimuli for which enhancement offers more limited benefit. Cancellation meshes well with the concept that perception involves a quest for invariance to irrelevant dimensions.

#### Cancellation as a Model of Sound.

The ability to cancel unwanted sounds is clearly useful for perception, but one might take a step further and argue that it is, in part, *constitutive* of perception. As a predictive model, a harmonic cancellation filter characterizes the part

of input that it can cancel, just as an autoregressive model characterizes its spectral envelope, or a binaural EC model its spatial position. The residual, which by definition does not fit that model, informs us about "what else is out there". It too can be characterized by recursively applying the same model or, alternatively, a compound model can be applied to the original sound to estimate parameters jointly (as in the multiple  $F_0$  model described in the Appendix, *Period Estimation*). This is related to concepts of predictive coding (Friston, 2018) and compression (Schmidhuber, 2009).

Like pattern classification (Duda et al., 2012), cancellation seeks invariance with respect to irrelevant dimensions of the input, specifically those that reflect the background. In contrast to classifiers that involve non-linear transforms, cancellation as described here is purely linear, which makes sense given that the acoustic mixing process itself is linear.

### How Useful is it in Practice?

Auditory Scene Analysis benefits from multiple cues and regularities, of which harmonicity is but one. Harmonic cancellation is likely to be useful in situations where neither temporal separation, nor spectral separation, nor binaural disparities are effective to suppress interfering sources, and then only if the interference is harmonic. Thus, at best, it is one tool among many, beneficial in a restricted set of circumstances.

Measured in terms of TMR at threshold performance, the harmonicity benefit can reach  $\sim$ 17 dB for identifying synthetic vowels, although most studies report smaller effects (Culling et al., 1994; de Cheveigné, Kawahara, et al., 1997; Summerfield, Culling, & Fourcin, 1992). This is of the same order of magnitude as reported for binaural unmasking (Colburn & Durlach, 1965; Jelfs, Culling, & Lavandier, 2011). In terms of proportion of tokens recognized, the benefit appears maximal for TMR around -15 dB and vanishes below -30 dB or above +15 dB

(de Cheveigné, 1999b; de Cheveigné, Kawahara, et al., 1997; McKeown, 1992). Thanks in part to harmonicity-based segregation, a target (wide-band harmonic or noise) mixed with a harmonic background can be detected at TMRs down to  $\sim -20$  dB (Gockel et al., 2002; Micheyl et al., 2006), or -32 dB for a narrowband noise target (Deroche & Culling, 2011a). The benefit relative to a noise or inharmonic masker is on the order of 5 to 15 dB (Deroche & Culling, 2011a; Deroche, Culling, & Chatterjee, 2014; Micheyl et al., 2006). Overall, harmonic cancellation mainly benefits *weak* targets.

For vowel identification, the benefit is measurable for  $\Delta F_0$  s as small as 0.4% but not less (de Cheveigné, 1997b), and plateaus for  $\Delta F_0$  s beyond  $\sim$ 6%. It is greater for longer stimuli (200 ms) than shorter stimuli (50 ms) (Assmann & Summerfield, 1994), but measurable for stimuli as short as four cycles of the lower  $F_0$  (23 ms at 175 Hz, McKeown & Patterson, 1995). It is reduced but not abolished if the masker's  $F_0$  is modulated at rates as fast as 5 Hz (200 ms period) (de Cheveigné, 1997b; Deroche & Culling, 2011b; Summerfield et al., 1992), suggesting a remarkable ability to track  $F_0$  variations. However, this breaks down in the presence of reverberation, whereas a similar degradation is not observed if the masker  $F_0$  is steady-state (Culling et al., 1994; Sayles, Stasiak, & Winter, 2015). Data from mistuned harmonic experiments suggest that the benefit might be limited to the spectral region below  $\sim$ 2-3 kHz (Hartmann et al., 1990). Indeed, in concurrent vowel experiments the benefit appears to stem mainly from the region below 1 kHz that includes a vowel's first formant (Culling & Darwin, 1993).

Real speech maskers differ from ideal harmonic maskers in that periodic portions are sparsely distributed over time (Hu & Wang, 2008), the  $F_0$  varies due to intonation, and periodicity is further degraded by articulation, irregularities in voice excitation, and added noise including reverberation. The benefit of a  $\Delta F_0$  between a monotonized speech target and monotonized masker (two concurrent

voices with the same  $F_0$ , or harmonic complex with spectral envelope similar to speech) ranges from 3-8 dB (Deroche & Culling, 2013; Deroche, Culling, Chatterjee, & Limb, 2014a; Deroche, Culling, Lavandier, & Gracco, 2017), which is also on the same order as binaural effects for similar stimuli (Deroche et al., 2017).

### Learning?

Pattern-matching models of pitch perception (de Boer, 1976) postulate some form of harmonic template, or "sieve" (Duifhuis, Willems, & Sluyter, 1982; Schroeder, 1968), and the same template is also required for a spectral domain model of segregation. This is non-trivial: the dictionary of templates must cover the full range of  $F_0$ s, there must be some mechanism to align the templates accurately with the substrate of frequency analysis (e.g. cochlea), and each template itself is a complex affair involving multiple slots with accurate tuning. It has been proposed that templates are learned from exposure to harmonic sounds such as speech (Bowling & Purves, 2015; Divenyi, 1979; Saddler, Gonzalez, & McDermott, 2020; Terhardt, 1974) possibly modulated by cultural preferences (J. McDermott & Hauser, 2004; J. H. McDermott, Lehr, & Oxenham, 2010; J. H. McDermott, Schultz, Undurraga, & Godoy, 2016; McPherson, Dolan, et al., 2020). The demonstration that templates can be learned from noise (Shamma & Dutta, 2019; Shamma & Klein, 2000) makes that argument more tenuous, and highlights the question of what, exactly, is being learned. Perhaps that algorithm discovers, rather than learns, the mathematical property that is exploited more directly by the cancellation filter.

The template-like properties of a time-domain cancellation filter (Eq. 1, Fig. 4) stem from mathematics, rather than learning. This is a big appeal: why jump through hoops when a simple solution is at hand? The organism may still need to discover that this regularity exists and is worth attending to, and the mechanism may need tuning, particularly if it involves combining frequency channels. This leaves ample room for learning, and possibly even cultural influences.

#### Is There Time?

In a classic chapter, de Boer (1976) likened auditory theory to a pendulum moving between "time" and "place" (spectrum). The pendulum is still swinging, and several recent papers have strengthened the case for spectral and placerate accounts (e.g. Shera et al., 2002; Su & Delgutte, 2020; Sumner et al., 2018; Verschooten, Desloovere, & Joris, 2018; Whiteford, Kreft, & Oxenham, 2020). Arguments for time remain (a) evidence for temporal mechanisms of binaural processing (see section *Analogy with Binaural Equalization-Cancellation* of the Appendix), (b) existence of specialized neural circuitry within the brain (see section *Anatomy and Physiology* of the Appendix), (c) the simplicity, effectiveness and ease of implementation of a time-domain harmonic filter, in contrast to a harmonic template or sieve in the frequency domain.

Hybrid models offer the best of both worlds, but they may worry scholars who care about parsimony or falsifiability. As a case in point, if we admit that delay might arise by cross-channel interaction (de Cheveigné & Pressnitzer, 2006), it is hard to say anything for, or against, the hypothesis that processing involves neural delays. On the other hand, it would be unwise to let this blind us to the possibility that that auditory system does rely on a combination of spectral and time-domain analysis.

My personal inclination is that auditory perception involves time-domain processing within the brain, but the effectiveness of that processing is enhanced by the peripheral bandpass filter bank that helps overcome the effects of nonlinear transduction and noise (based on principles related to Logan's theorem). High-resolution mechanical filtering serves to "pre-calculate" a set of useful basis functions upon which the brain then operates in the time-domain (see sections *Transforms in Filter Space* and *Non-Linearity* of the Appendix). In this perspective, cochlear mechanics are the "last chance" to process acoustical signals with good

resolution, linearity, and low noise, before handing transduced patterns over to more flexible but less accurate neural processing.

Carving Sound at its Joints. Auditory Scene Analysis is often described as a process of assembling elements across the spectrum (simultaneous grouping) or across time (sequential grouping) (Bregman, 1990), mirroring the common process of additive or concatenative synthesis by which stimuli are created in the lab. It glosses over the issue of whether these ingredients are recoverable from the mix, upon which this assumption depends. Once the coins are thrown into the melting pot, can we pull them out intact? According to classic Auditory Scene Analysis, we can: spectral analysis reveals "natural kinds" (partials), between which are found the "joints" at which sounds may be carved (Campbell, O'Rourke, & Slater, 2011). Indeed, according to this view, a grouping mechanism is required for any complex sound to form a coherent whole, otherwise it might shatter into as many percepts as partials (even though few of us would claim to ever have heard more than a couple of such percepts within a sound). The wisdom of invoking sinusoidal partials as "natural kinds" on which Auditory Scene Analysis processes operate is rarely questioned.

In contrast, harmonic cancellation requires no analysis-into-parts or grouping. Whereas a bandpass filter is defined by what it selects (a frequency band), a cancellation filter is defined by what it removes (periodic power at period T). This is an example, like a shadow, of what Sorensen (2011) calls a "para-natural kind". The process is effective both to characterize a periodic sound by its parameter T, and to get rid of that sound and search for more. It is an alternative way to "carve sound at its joints".

# **Conclusion**

The harmonic cancellation hypothesis states that the harmonic (or periodic) structure of interfering sounds can be exploited to suppress or ignore them. A large body of experimental results are consistent with this hypothesis, whereas alternative hypotheses for  $F_0$ -based segregation are less well supported. In particular, harmonic enhancement, according to which harmonicity of a target makes it resilient to masking, receives little support, which is surprising because counter to our intuition and inconsistent with textbook explanations of scene analysis involving a harmonicity-based "grouping" operation. Harmonic cancellation fits well with an account of perception as seeking invariance with respect to irrelevant dimensions of the sensory pattern, and with the concept of "unconscious inference" promoted by Helmholtz. Harmonic cancellation can be implemented in the frequency domain (based on cochlear analysis) or time domain (based on the temporal processing of neural discharge patterns). Support for the latter comes from the success of the related EC model of binaural interactions, from the presence of neural structures apparently specialized for processing of temporal information, and from theoretical considerations that suggest that a time-domain implementation might be more straightforward and effective.

# **Appendix: Deeper Issues**

The harmonic cancellation hypothesis is straightforward and well supported experimentally, but it raises a number of interesting questions that are worth considering.

#### Hybrid models

The hybrid harmonic cancellation models enumerated in the main text are described here in greater detail.

• Hybrid model 1: Cancellation-enhanced spectral patterns. Each channel of a filter bank is convolved with a cancellation filter tuned to T. This has the effect of sharpening spectral analysis so that the outcome is closer to the ideal (Fig. 2 right). The pattern of power over channels is then handed over to a frequency-domain pattern-matching stage. This is illustrated in Fig. 6 (a). Two vowels, /a/ and /e/ with fundamentals 100 Hz and 106 Hz respectively (left), are mixed. Cues to /e/ are indistinct within the spectrum of the mix (right, black), but can be enhanced by applying to each channel a cancellation filter tuned to suppress /a/ (right, red). This model is reminiscent of periodicity tagging of tonotopic patterns (Keilson, Richards, Wyman, & Young, 1997), or of the place-time model of Assmann and Summerfield (1990) in which a spectral profile for the target vowel was taken by sampling the autocorrelation function at the target's period. If the spectral profile were derived from a limited window of cancellation-filtered signal, placing that window within the background-invariant part (red in Fig. 4 (b), right) would make the profile invariant with respect to backgrounds of period T. The pattern would still be distorted by the cancellation filtering, and spectral pattern-matching would need to take this into account.

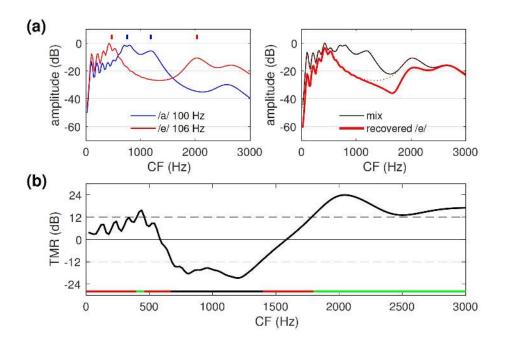


Figure 6: Two hybrid models of harmonic cancellation. (a) Hybrid Model 1. Left: power as a function of CF for synthetic vowels /a/,  $F_0$ =100 Hz (blue) and /e/,  $F_0$ =106 Hz (red). Short lines above the plot indicate the first two formant frequencies of each vowel. Right: power as a function of CF for the mix before (black) and after (red) applying a cancellation filter tuned to suppress the period of /a/. (b) Hybrid Model 3. Black: per-channel TMR of vowel /e/ as a function of CF for a mixture of /a/+/e/ at overall TMR=0 dB. Channels are divided into 3 groups: TMR>12 dB (green, to be left intact), TMR<-12 dB (black, to be discarded) and -12 dB $\leq TMR\leq 12$  dB (red, to be filtered by a cancellation filter).

• Hybrid Model 2: Channel rejection on the basis of periodicity. Filter bank channels are divided into two groups based on TMR (estimated based on residual power at the output of a cancellation filter tuned to T). The first group consists of channels dominated by the background; these are rejected. The remaining channels are handed over to the pattern-matching stage to be matched based on their temporal pattern. This principle was employed in the concurrent vowel identification model of Meddis and Hewitt (1992), itself inspired from earlier ideas for binaural or periodicity-based segregation (Lyon, 1983, 1988; Weintraub, 1985). Spectral resolution must be sufficient

so that enough channels are spared to represent the target.

• Hybrid Model 3: Cancellation filtering of selected channels. Filter bank channels are divided into three groups based on TMR. Channels with large TMR are left untouched, channels with small TMR are discarded, and intermediate channels are processed by the cancellation filter. Keeping the first group intact reduces target distortion, and discarding the second group avoids contamination from noise if the cancellation filter is imperfect (as it might be due to nonlinearity or noise). Cancellation filtering is reserved for channels with intermediate TMR, for which it can be effective. This model differs from Hybrid model 2 by the presence of this third group. A similar suggestion was made by Guest and Oxenham (2019).

Hybrid Model 3 is illustrated in Fig. 6 (b). The black line shows the TMR per channel at the output of a filter bank in response to the mix /a/+/e/ with overall TMR = 0 dB. Channels for which TMR exceeds some threshold (+12 dB in this example) are left intact (green), channels for which TMR is below a second threshold (-12 dB in this example) are discarded (black). Channels with intermediate TMR are processed with a cancellation filter (red).

- Hybrid Model 4: Channel-specific cancellation filter. In contrast to previous models, for which the parameter T is the same for all channels, here it is allowed to vary across channels. This is analogous to the channel-dependent versions of the EC model of binaural unmasking (Akeroyd, 2004; Breebaart et al., 2001; Culling & Summerfield, 1994). This hypothesis may be useful to explain results found with inharmonic stimuli (e.g. Roberts & Brunstrom, 1998) as discussed in the main text.
- Hybrid Model 5: Synthetic delays. The cancellation filter of Eq. 1 re-

quires a delay equal to the background period (e.g. 20 ms for a 50 Hz fundamental). The existence of delays of this size in the auditory system has been questioned (e.g. Laudanski, Zheng, & Brette, 2014), and to address this issue it has been suggested that long delays might arise from cross-channel interaction (de Cheveigné & Pressnitzer, 2006). According to this model, the filter bank serves mainly that purpose: to help synthesize the delay T required by Eq. 1.

• Hybrid Model 6: Logan's theorem. Rather than a specific model, this is a processing principle that addresses the issue of the non-linear transduction that follows cochlear filtering. Due to half-wave rectification, each transduced signal is "blind" to one-half of every cycle, and thus one might worry that some information was lost. Logan's theorem states instead that a narrowband signal can be reconstructed perfectly from its zero crossings, and hence also from its half-wave rectified version (Logan, 1977; Shamma & Lorenzi, 2013). To the extent that it is applicable here, the benefit of cochlear filtering would be to *linearize* transduction, so that neural signal processing has, in effect, full access to the acoustic waveform (see the section *Non-Linearity* below).

#### Period Estimation

Harmonic cancellation requires an estimate of the interferer period T. Harmonic cancellation itself can be used for that purpose: an array of cancellation filters, each tuned to a different delay (lag) covering the range of expected periods, shows a minimum in output power at a lag equal to the period. This is equivalent to searching for a peak in the autocorrelation function (de Cheveigné, 1998; Licklider, 1951; Meddis & Hewitt, 1991). The relation between cancellation and correlation is detailed in the next section.

From this perspective, cancellation is both an *analysis* tool (it cancels part of a signal to reveal the remainder), and an *estimation* tool (it estimates the period of the part it cancels). Applied recursively to a mixture of two sounds, it can reveal two periods: we first estimate the period of the dominant sound and cancel it, and then recurse on the remainder. These steps can be performed in parallel by searching the two-dimensional parameter space of a cascade of cancellation filters defined as  $h_1(t) = \delta_0(t) - \delta_{\tau_1}(t)$  and  $h_2(t) = \delta_0(t) - \delta_{\tau_2}(t)$  for a minimum in output power. This output is zero when  $[\tau_1, \tau_2] = [nT_1, mT_2]$  for integers m, n (de Cheveigné, 1993; de Cheveigné & Kawahara, 1999). Interestingly, a neural version of this model designed to estimate the pitch of a mistuned partial (de Cheveigné, 1999a) accurately accounted for the subtle shifts observed by Hartmann and Doty (1996); Hartmann et al. (1990), see also Holmes and Roberts (2012)

Associated with the period is an estimate of the degree to which the sound is, in fact, periodic. A straightforward measure is output power of a cancellation filter tuned to the period T, normalized by power at the input (or by output averaged over other lags, e.g.  $1 \dots T$ ). A value of zero indicates that the sound is perfectly periodic, and a small value indicates that it is "approximately periodic". This same measure can be used as a criterion to detect a target in the presence of a harmonic background.

The threshold beyond which a sound should be declared "aperiodic" depends on the application, and more specifically on the distributions of "periodic" and "aperiodic" sounds as defined by the application's needs. It is worth noting that residual aperiodic power at the output of a narrowband filter (e.g. filter bank channel) takes on relatively low values even if the stimulus is aperiodic. The threshold needs adjusting accordingly.

#### Correlation and Cancellation

We can define the running autocorrelation function (ACF) at time t as

$$r_t(\tau) = \sum_{i=t}^{t+W} s(i)s(i-\tau)$$
(3)

(dropping the scaling factor 1/W for simplicity), where W is the duration of a sliding integration window that serves to smooth the time course of  $r_t$ . Power at time t can then be defined as  $P_t = r_t(0)$ . Likewise, we can define a squared difference function (SDF) as power at time t of the cancellation filter output:

$$d_t(\tau) = \sum_{i=t}^{t+W} [s(i) - s(i-\tau)]^2.$$
 (4)

ACF and SDF are then related by:

$$2r_t(\tau) = P_t + P_{t-\tau} - d_t(\tau).$$
 (5)

A peak in correlation, cue to the period, maps to a trough in difference function. It is convenient to normalize ACF and SDF:

$$\bar{r}_t(\tau) = r_t(\tau)/\sqrt{P_t P_{t-\tau}},$$
 (6)

$$\bar{d}_t(\tau) = \sum_{i=t}^{t+W} \left[ s(i) / \sqrt{P_t} - s(i-\tau) / \sqrt{P_{t-\tau}} \right]^2,$$
 (7)

in which case the normalized functions are related more simply by

$$2\bar{r}_t(\tau) = 1 - \bar{d}_t(\tau). \tag{8}$$

For a periodic sound with period T,  $\bar{r}_t(T) = 1$ , and  $\bar{d}_t(T) = 0$ .

Equation 5 is useful to derive the ACF from the SDF or vice-versa. It can also be extended to more terms, for example to implement a cascade of cancel-

lation filters in terms of correlation. This allows different modeling strands to be unified, and justifies some flexibility when speculating about hypothetical neural implementations (see below).

#### Analogy with Binaural Equalization-Cancellation

Durlach's EC model has been successful in accounting for binaural unmasking (Culling, 2007; Culling & Summerfield, 1994; Durlach, 1963) and binaural pitch phenomena (Culling, Summerfield, & Marshall, 1998), and in predictive models of speech intelligibility (Beutelmann & Brand, 2006; Cosentino, Marquardt, McAlpine, Culling, & Falk, 2014; Lavandier et al., 2012; Schoenmaker, Brand, & van de Par, 2016). Binaural interaction has also been couched in terms of inter-aural *correlation* rather than cancellation (Jeffress, 1948) but, as pointed out by D. M. Green (1992), an EC model can be implemented on the basis of interaural correlation, and vice versa, as the two are related:  $[s_L(t) - \alpha s_R(t-\tau)]^2 = s_L(t)^2 + \alpha^2 s_R(t-\tau)^2 - 2\alpha s_L(t) s_R(t-\tau)$ , where  $s_L$  and  $s_R$  are sounds at left and right ears, respectively. A cancellation residue in one model maps to decorrelation in the other.

An interesting suggestion is that EC might operate independently within frequency channels (Akeroyd, 2004; Breebaart et al., 2001; Culling & Summerfield, 1994), rather than with parameters common to all channels as in the original EC model (Durlach, 1963). It has been further suggested that EC parameters can be estimated and applied within short time windows (Hauth & Brand, 2018; Wan, Durlach, & Colburn, 2014), which paves the way for a spectro-temporal form of the EC model that supports "glimpsing" (Beutelmann, Brand, & Kollmeier, 2010).

A monaural version of the EC model has been invoked to explain comodulation masking release (CMR) (Piechowiak, Ewert, & Dau, 2007).

Anatomy and Physiology

Time-domain and hybrid models entail time-domain signal processing within the brain. Anatomical and physiological specializations to support such processing include transduction and coding of acoustic temporal structure in the auditory nerve (up to 4-5 kHz or possibly higher, Carcagno, Lakhani, & Plack, 2019; Hartmann, Cariani, & Colburn, 2019; Heinz, Colburn, & Carney, 2001; Verschooten et al., 2019), specialized synapses in the cochlear nucleus and subsequent relays, and fast excitatory and inhibitory interaction in the medial and lateral superior olives (MSO and LSO) (Beiderbeck et al., 2018; Grothe, 2000; Keine, Rübsamen, & Englitz, 2016; Stasiak, Sayles, & Winter, 2018; Zheng & Escabí, 2013) and other nuclei (Albrecht, Dondzillo, Mayer, Thompson, & Klug, 2014; Caspari, Baumann, Garcia-Pino, & Koch, 2015; Felix et al., 2017). Some of these circuits are interpreted as serving binaural interaction, but presumably could be borrowed for other needs (see Joris & van der Heijden, 2019; Kandler, Lee, & Pecka, 2020, for recent reviews).

The time-domain cancellation filter of Fig. 4 (c, left), Eq. 1, can be approximated by the "neural cancellation filter" of Fig. 4 (c, right). Spikes arriving via the direct pathway are suppressed by the coincident arrival of spikes delayed by T. Applied to data recorded from the auditory nerve in response to a mixture of two vowels with different  $F_0$ s (Palmer, 1990), that simple circuit was effective in estimating both their periods and suppressing correlates of one or the other vowel (de Cheveigné, 1993, 1997a; Guest & Oxenham, 2019). Such a mechanism would require temporally-accurate neural representations (excitatory and inhibitory), delays, and an inhibitory gating or "anticoincidence" mechanism.

Temporally-accurate inhibitory transforms of sensory input are created in several nuclei, including cochlear nucleus (CN) (stellate-D cells), medial and lateral nuclei of trapezoid body (MNTB and LNTB), and ventral nucleus of the lateral lemniscus (VNLL) (Arnott, Wallace, Shackleton, & Palmer, 2004; Caspari et al.,

2015; Joris & Trussell, 2018). Fast interaction between direct and delayed neural patterns could in principle occur as early as the dendritic fields of cells in CN (Davis & Voigt, 1997; Needham & Paolini, 2006; Schofield, 1994; Shore, Helfert, Bledsoe, Altschuler, & Godfrey, 1991; Xie & Manis, 2013), or as late as dendritic fields of the inferior colliculus (IC) (Caspari et al., 2015; Chen, Read, & Escabí, 2019). A recent study reported evidence for an inhibitory "veto" mechanism at the axon initial segment of LSO principal neurons, with very narrow tuning to inter-aural time differences (Franken et al., 2021). Transmission failure at reputed "secure" synapses in CN and MNTB might conceivably reflect a similar veto mechanism (Englitz, Tolnai, Typlt, Jost, & Rübsamen, 2009; Mc Laughlin, van der Heijden, & Joris, 2008; Stasiak et al., 2018).

The cancellation-correlation equivalence discussed earlier implies that fast interaction might also be excitatory-excitatory, the correlation pattern being later converted to a cancellation-like statistic by slower inhibitory interaction along the lines of Eqs. 5 and 8. Note, however, that finding a minimum of cancellation would then require subtraction of two large correlation values, which may be a problem if those values are coded by a representation (like rate of a Poisson-like process) for which the noise variance of the value increases with its mean. One might speculate that the cost of specialized fast inhibitory circuitry is recouped by the benefit of performing cancellation directly.

There is also evidence in favor of accurate rate-place spectral representations (Fishman, Micheyl, & Steinschneider, 2013; Fishman, Steinschneider, & Micheyl, 2014; Larsen, Cedolin, & Delgutte, 2008; Su & Delgutte, 2020) that might support a spectral version of the harmonic cancellation hypothesis, particularly as it has been argued that tuning might be narrower in humans than in most model animals (Shera et al., 2002; Sumner et al., 2018; Verschooten et al., 2018; Walker, Gonzalez, Kang, McDermott, & King, 2019). Narrow tuning might also benefit a

spectro-temporal mechanism, with the caveat that narrower filters are temporally more sluggish.

Sinex et al. (2002); Sinex and Li (2007); Sinex, Li, and Velenovsky (2005) report stronger responses in IC neurons for mistuned partials, consistent with the output of a cancellation filter, but they explain it by a different model based on cross-channel interaction of between-partial beat patterns, analogous to the waveform interaction models described earlier. Their model also accounts for the particular temporal structure of the response; whether that structure too could be explained by cancellation remains to be determined.

In summary, known neural circuitry might support both temporal and spectral mechanisms of harmonic cancellation, however I am not aware of evidence as strong as that reported in favor of the EC model. A rate-frequency response such as Fig. 4 (a) might evade notice if attention is devoted to *peaks* of activity rather than dips. It could also elude discovery if the output pattern follows a latency code rather than rate code (Chase & Young, 2007). The filter output in Fig. 4 (b) is evocative of ON-OFF patterns observed in the superior paraolivary nucleus (SPON) (Kandler et al., 2020) but this similarity could be fortuitous, indeed those patterns have been attributed to gap detection or duration encoding (Kadner & Berrebi, 2008).

#### Smart Pattern Matching

As discussed in the main text (*Harmonic Cancellation – Possible Mechanisms*), each recovered target pattern is affected by two error terms: imperfect cancellation of the background, and distortion undergone by the target. In the time-domain model, the first term can be reduced to zero over part of the pattern (red segment in Fig. 4 b, right). This assumes the ability to locate and isolate reliable intervals, which is commonly granted for auditory perception (Moore et al., 1988; Viemeister & Wakefield, 1991).

There remains the second error term due to filter-induced target distortion. This can be mitigated *if it is known to the pattern matching stage*, for example by applying the same distortion to each pattern in the dictionary. Distortion consists of an attenuation factor applied to each target component depending on how close it falls to the harmonic series of the background, as quantified by the filter transfer function (Fig. 4 a, right). This produces a "moiré effect" that can be quantified (and thus taken into account) if  $F_0$ s of both *background and target* are known.

Target patterns can be further refined if the background is stationary over more than two periods, as illustrated in Fig. 7. Specifically, if the stimulus is long enough to define N distinct observation intervals temporally separated by T, these intervals can form N(N-1)/2 distinct pairs from which to infer the target. These observations are not all strictly independent, but the distortion (Fig. 7 right) and noise patterns differ between pairs and this may assist inference. A perceptual mechanism operating in this fashion might seem implausibly complex. On the other hand, we cannot rule out that the trick is discovered by a learning process. The point made here is that the opportunity exists.

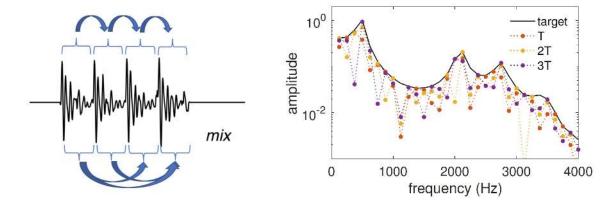


Figure 7: Left: waveform of the mix of target vowel /e/ (132 Hz) with background vowel /a/ (100 Hz) at TMR=-12 dB. Given four background cycles, intervals can be paired over spans of T, 2T and 3T, with three, two and one repeats respectively (blue arrows). Right: spectrum of target vowel /e/ (black line) and cancellation-filtered estimates obtained for spans T, 2T and 3T (symbols). Averaging over estimates (or better: taking their maximum) would yield a more accurate estimate of the target, and averaging over repeats might further attenuate uncorrelated noise (not shown).

### Transforms in Filter Space

The idea that *cochlear filtering* works hand in hand with *neural filtering* is intriguing. What are the possibilities, what are the limits? As an example, the bandwidth of cochlear filters is usually seen as a hard limit on spectral resolution, but it appears that with neural filtering that limit can be overcome, as exploited by past schemes such as the "second filter" (Huggins & Licklider, 1951), stereausis (Shamma, Shen, & Gopalaswamy, 1989), lateral inhibitory network (LIN) (Shamma, 1985), phase opponency (Carney, Heinz, Evilsizer, Gilkey, & Colburn, 2002), synthetic delays (de Cheveigné & Pressnitzer, 2006), EC (Durlach, 1963), selectivity focusing in inferior colliculus (IC) (Chen et al., 2019), and here harmonic cancellation.

This section attempts to make sense of this situation by casting both filtering stages into a common framework. Any filter can be approximated as a fi-

nite impulse response filter (FIR) of order N, defined by the column vector  $\mathbf{h} = [h_0, h_1, \dots, h_N]^{\top}$  of impulse response coefficients. A signal s(t) is filtered by convolving it with this impulse response. Alternatively, using matrix notation, if  $\mathbf{S} = [s(t), s(t-1), \dots s(t-N+1)]$  is the  $T \times N$  matrix of time-lagged signals, the filtered signal is obtained as the product  $\mathbf{Sh}$ . A useful way to think of it is that the lags  $[0 \dots N]$  create a *memory* of the past signal, within which the filter can "shop" for useful information to characterize variations over time.

Extending to a M-channel filter bank, the filters can be defined by a matrix of impulse responses  $\mathbf{F}$  of size  $N \times M$ , where each column of  $\mathbf{F}$  represents the impulse response of one channel. The matrix of filtered signals is then obtained as the product  $\mathbf{S}' = \mathbf{S}\mathbf{F}$ . To relate this to the context of this paper, picture s(t) as an acoustic signal,  $\mathbf{F}$  as a bank of "cochlear" filters, and  $\mathbf{S}'$  as a matrix of vibration waveforms at different points along the basilar membrane.

If the matrix  $\mathbf{F}$  is of rank N, it has a right inverse  $\mathbf{\bar{F}}$  such that  $\mathbf{F}\mathbf{\bar{F}}=\mathbf{I}$ , the identity matrix. Why might this be useful? Suppose that we wish to speculate that the auditory brain implements a particular filter (defined by its impulse response h applicable to the acoustic waveform). We know that it does not have access to time-lagged acoustic signals  $\mathbf{S}$ , so it cannot implement that filter directly, but it does have access to peripheral filter outputs  $\mathbf{S}'$ . We want to know if our speculation is realistic.

We can write:

$$\mathbf{Sh} = \mathbf{SF\bar{F}h} = \mathbf{S}'(\bar{\mathbf{F}h}) = \mathbf{S}'\mathbf{h}',\tag{9}$$

where  $\mathbf{h}' = \mathbf{\bar{F}}\mathbf{h}$  is a vector of weights. Applying weights  $\mathbf{h}'$  to  $\mathbf{S}'$  yields the desired filtered signal, exactly as if we had applied the filter  $\mathbf{h}$  directly to the acoustic waveform. Whereas the filter was originally defined by its coordinates  $\mathbf{h}$  on a basis of time shifts applicable to the acoustic signal, it is now defined using

coordinates h' on a basis of filter bank channels. The outcome is the same.

Why is this relevant here? It means that essentially any filter can be implemented (or its implementation can be complemented) by forming a weighted sum of cochlear filter outputs, as long as their impulse responses are long enough to reach the required order N. This is the gist of the "synthetic delay" model of de Cheveigné and Pressnitzer (2006). According to this view, peripheral filtering and neural time-domain interaction work hand in hand to perform acoustic signal processing (subject to limits imposed by noise and nonlinearity discussed in the next section).

A matrix of N cancellation filters with lag parameters T ranging from 0 to N-1 is also invertible (if one replaces the degenerate T=0 filter by  $\delta_0(t)$ ), and thus one can treat it as a "basis" similar to the filter bank basis just described. A filter defined by its coefficients  $\mathbf{h}$  on a lags basis, or  $\mathbf{h}'$  on a filter bank basis, can therefore also be defined by a set of coefficients  $\mathbf{h}''$  on this new basis. One can, at least conceptually, transform the sensory representation back and forth between these three representations: lagged waveforms, band pass filter bank channels, and cancellation-filtered channels, with no loss of information. The cancellation-filtered representation is reminiscent of the pitch-like "level of representation" invoked by Hafter and Saberi (2001).

There remains one difficult issue: given a periodic sound with period T, how do we find the coefficients  $\mathbf{h}'$  of a cancellation filter (defined over a basis of peripheral filter outputs) that can cancel it? In the standard formulation (Eq. 1) based on a basis of lags, the filter  $\mathbf{h}$  consists of all zeros except h(0)=1 and h(T)=-1, so the parameter T can easily be found by scanning a linear array for a minimum. For  $\mathbf{h}'$ , the situation is more complex because we must find a set of N parameters, rather than one, to obtain the same result. This is a serious obstacle unless a "smart" way of finding  $\mathbf{h}'$  is found. A full discussion of the problem is beyond the

scope of this paper, but it is worth taking note of three points.

The first is that, if principal component analysis (PCA) is applied to the matrix S for a periodic input with period  $T \leq N$ , at least one column of the PCA transform matrix defines a FIR filter S that cancels that input. This is because the S the column of S is identical to the 0th column (periodicity), hence S is not of full rank.

The second point follows from the first: if PCA is applied to the matrix S' of filter bank outputs, at least one column of the PCA transform matrix defines a set of coefficients h' that also cancels its input. This is because rank deficiency of S implies rank-deficiency of S'. Thus, the appropriate coefficients h' can be also be found by applying PCA to filter bank outputs for a periodic input. This data-dependent process can be seen as a form of data-driven learning, analogous to what we discussed earlier.

The third point is that PCA is widely considered as a plausible neural operation (Minden, Pehlevan, & Chklovskii, 2018; Oja, 1982; Qiu, Wang, Lu, Zhang, & Du, 2012). Putting these pieces together, we can speculate that the hypothesis that Eq. 1 is implemented in the brain as a weighted sum of filter bank outputs, rather than a simple delay T, is not completely unrealistic. This rough sketch needs fleshing out, but it suggests a possible direction to model how the auditory brain might implement complex signal processing tasks, cancellation being one particular example.

Again, such operations might seem implausibly complex for a biological implementation, but knowing that the option exists, in principle, and understanding how it works, can guide speculation that something similar is discoverable by a learning process.

#### Non-Linearity

Previous sections mostly glossed over the issue of non-linear transduction.

The suggestion that linear operations can be swapped, as in Fig. 5 (b), or linear transforms inverted as in the previous paragraph, is moot if the systems are not linear. What can be salvaged from those simple ideas?

First, note that any time-invariant transform of a periodic signal is periodic with the same period (or submultiple of that period), so a cancellation filter tuned to the period would produce zero output as in the linear case. Thus, for example, Hybrid Model 1 would work as advertized. Second, pattern distortions due to nonlinearity may be compensated for in the pattern-matching stage. Thus, Hybrid Model 2 might also work. Third, more generally, we can invoke Logan's theorem and assume that the deleterious effects of nonlinearity, whatever they are, can be redressed by subsequent processing. The theorem doesn't say how, but it is easy to imagine simple situations in which this might pan out. For example, sampling the steep phase characteristic of the cochlear filter bank at two points differing by  $\pi$  might give access to both polarities of the signal at that point, reversing effects of half-wave rectification. Fourth, non-linearity demodulates the band-pass filtered signal, thus abstracting an informative temporal envelope from less robust fine structure (Dau et al., 1997). In this respect, non-linearity is a feature, rather than a bug.

In summary, non-linearity does not prevent harmonic cancellation, although it does make it harder to understand the limits of what can be achieved, and how.

## **References**

- Akeroyd, M. A. (2004). The across frequency independence of equalization of interaural time delay in the equalization-cancellation model of binaural unmasking. *J. Acoust. Soc. Am.*, 116, 1135-1148. doi: 10.1121/1.1768959
- Albrecht, O., Dondzillo, A., Mayer, F., Thompson, J. A., & Klug, A. (2014). Inhibitory projections from the ventral nucleus of the trapezoid body to the medial nucleus of the trapezoid body in the mouse. *Frontiers in Neural Circuits*, 8. doi: 10.3389/fncir.2014.00083
- al Haytham, I. (1030). Book of optics (in Hatfield, 2002).
- Arehart, K. H., Rossi-Katz, J., & Swensson-Prutsman, J. (2005). Double-Vowel Perception in Listeners With Cochlear Hearing Loss: Differences in Fundamental Frequency, Ear of Presentation, and Relative Amplitude. *Journal of Speech, Language, and Hearing Research*, 48, 236–252. doi: 10.1044/1092-4388(2005/017)
- Arehart, K. H., Souza, P. E., Muralimanohar, R. K., & Miller, C. W. (2011). Effects of Age on Concurrent Vowel Perception in Acoustic and Simulated Electroacoustic Hearing. *Journal of Speech, Language, and Hearing Research*, *54*, 190–210. doi: 10.1044/1092-4388(2010/09-0145)
- Arnott, R., Wallace, M., Shackleton, T., & Palmer, A. (2004). Onset Neurones in the Anteroventral Cochlear Nucleus Project to the Dorsal Cochlear Nucleus. *Journal of the Association for Research in Otolaryngology*, 5. doi: 10.1007/s10162-003-4036-8
- Assmann, P. F., & Summerfield, Q. (1990). Modeling the perception of concurrent vowels: Vowels with different fundamental frequencies. *J Acoust Soc Am*, 88, 680–697. doi: 10.1121/1.399772
- Assmann, P. F., & Summerfield, Q. (1994). The contribution of waveform interactions to the perception of concurrent vowels. *J Acoust Soc Am*, 95, 471–484. doi:10.1121/1.408342
- Beiderbeck, B., Myoga, M. H., Müller, N. I. C., Callan, A. R., Friauf, E., Grothe, B., & Pecka, M. (2018). Precisely timed inhibition facilitates action potential firing for spatial coding in the auditory brainstem. *Nature Communications*, *9*, 1771. doi: 10.1038/s41467-018-04210-y
- Bernstein, J. G. W., & Oxenham, A. J. (2008). Harmonic segregation through mistuning can improve fundamental frequency discrimination. *J Acoust Soc Am*, 124, 1653–1667. doi: 10.1121/1.2956484
- Best, V., Roverud, E., Baltzell, L., Rennies, J., & Lavandier, M. (2019). The importance of a broad bandwidth for understanding "glimpsed" speech. *J Acoust Soc Am*, *146*, 3215–3221. doi: 10.1121/1.5131651
- Beutelmann, R., & Brand, T. (2006). Prediction of speech intelligibility in spatial noise and reverberation for normal-hearing and hearing-impaired listeners.

- J Acoust Soc Am, 120, 331-342. doi: 10.1121/1.2202888
- Beutelmann, R., Brand, T., & Kollmeier, B. (2010). Revision, extension, and evaluation of a binaural speech intelligibility model. *J Acoust Soc Am*, 127, 2479–2497. doi: 10.1121/1.3295575
- Binns, C., & Culling, J. F. (2007). The role of fundamental frequency contours in the perception of speech against interfering speech. *J Acoust Soc Am*, 122, 1765–1776. doi: 10.1121/1.2751394
- Bowling, D. L., & Purves, D. (2015). A biological rationale for musical consonance. *Proceedings of the National Academy of Sciences*, 112, 11155–11160. doi: 10.1073/pnas.1505768112
- Breebaart, J., van de Par, S., & Kohlrausch, A. (2001). Binaural processing model based on contralateral inhibition. I. Model structure. *J Acoust Soc Am*, 110, 1074–1088. doi: 10.1121/1.1383297
- Bregman, A. S. (1990). Auditory scene analysis. Cambridge, Mass.: MIT Press.
- Brokx, J., & Nooteboom, S. (1982). Intonation and the perceptual separation of simultaneous voices. *Journal of Phonetics*, 10, 23–36. doi: 10.1016/S0095-4470(19)30909-X
- Campbell, J. K., O'Rourke, M., & Slater, M. H. (Eds.). (2011). *Carving nature at its joints: natural kinds in metaphysics and science*. Cambridge, Mass: MIT Press.
- Carcagno, S., Lakhani, S., & Plack, C. J. (2019). Consonance perception beyond the traditional existence region of pitch. *J Acoust Soc Am*, *146*, 2279–2290. doi: 10.1121/1.5127845
- Carney, L. H., Heinz, M. G., Evilsizer, M. E., Gilkey, R. H., & Colburn, H. S. (2002). Auditory Phase Opponency: A Temporal Model for Masked Detection at Low Frequencies. *Acta Acust. Acust.*, 88, 15.
- Carney, L. H., Li, T., & McDonough, J. M. (2015). Speech Coding in the Brain: Representation of Vowel Formants by Midbrain Neurons Tuned to Sound Fluctuations1,2,3. *eNeuro*, *e0004-15.2015 1X12*. doi: 10.1523/ENEURO .0004-15.2015
- Caspari, F., Baumann, V. J., Garcia-Pino, E., & Koch, U. (2015). Heterogeneity of Intrinsic and Synaptic Properties of Neurons in the Ventral and Dorsal Parts of the Ventral Nucleus of the Lateral Lemniscus. *Frontiers in Neural Circuits*, 9. doi:10.3389/fncir.2015.00074
- Chalikia, M. H., & Bregman, A. S. (1993). The perceptual segregation of simultaneous vowels with harmonic, shifted, or random components. *Perception & Psychophysics*, *53*(2), 125–133. doi: 10.3758/BF03211722
- Chase, S. M., & Young, E. D. (2007). First-spike latency information in single neurons increases when referenced to population onset. *Proceedings of the National Academy of Sciences*, 104(12), 5175–5180. doi: 10.1073/pnas.0610368104

- Chen, C., Read, H. L., & Escabí, M. A. (2019). A temporal integration mechanism enhances frequency selectivity of broadband inputs to inferior colliculus. *PLOS Biology*, *17*(6), e2005861. doi: 10.1371/journal.pbio.2005861
- Colburn, H. S., & Durlach, N. I. (1965). Time-Intensity Relations in Binaural Unmasking. *J Acoust Soc Am*, 38(1), 93–103. doi: 10.1121/1.1909625
- Cooke, M. (2006). A glimpsing model of speech perception in noise. *J Acoust Soc Am*, 119(3), 1562–1573. doi: 10.1121/1.2166600
- Cooke, M., Morris, A., & Green, P. (1997). Missing data techniques for robust speech recognition. In *ICASSP* (Vol. II, p. 863-866).
- Cosentino, S., Marquardt, T., McAlpine, D., Culling, J. F., & Falk, T. H. (2014). A model that predicts the binaural advantage to speech intelligibility from the mixed target and interferer signals. *J Acoust Soc Am*, *135*(2), 796–807. doi: 10.1121/1.4861239
- Culling, J. F. (2007). Evidence specifically favoring the equalization-cancellation theory of binaural unmasking. *J. Acoust. Soc. Am.*, 122(5), 2803-2813. doi: 10.1121/1.2785035
- Culling, J. F., & Darwin, C. J. (1993). Perceptual separation of simultaneous vowels: Within and across-formant grouping by  $F_0$ . JAcoust Soc Am, 93(6), 3454-3467. doi: 10.1121/1.405675
- Culling, J. F., & Darwin, C. J. (1994). Perceptual and computational separation of simultaneous vowels: Cues arising from low-frequency beating. *J Acoust Soc Am*, 95(3), 1559–1569. doi: 10.1121/1.408543
- Culling, J. F., Hodder, K. I., & Toh, C. Y. (2003). Effects of reverberation on perceptual segregation of competing voices. *J Acoust Soc Am*, 114(5), 2871. doi: 10.1121/1.1616922
- Culling, J. F., Summerfield, A. Q., & Marshall, D. H. (1998). Dichotic pitches as illusions of binaural unmasking. I. Huggins' pitch and the "binaural edge pitch". *J Acoust Soc Am*, 103(6), 3509–3526. doi: 10.1121/1.423059
- Culling, J. F., & Summerfield, Q. (1994). Binaural segregation of concurrent sounds involves within-channel rather than across-channel processes. *J Acoust Soc Am*, 95(5), 2915–2915. doi: 10.1121/1.409275
- Culling, J. F., Summerfield, Q., & Marshall, D. H. (1994). Effects of simulated reverberation on the use of binaural cues and fundamental-frequency differences for separating concurrent vowels. *Speech Communication*, *14*, 71–95. doi: 10.1016/0167-6393(94)90058-2
- Darwin, C. J., & Bethell-Fox, C. E. (1977). Pitch Continuity and Speech Source Attribution. *Journal of Experimental Psychology: Human Perception and Performance*, *3*, 665–672. doi: 10.1037/0096-1523.3.4.665
- Dau, T., Kollmeier, B., & Kohlrausch, A. (1997). Modeling auditory processing of amplitude modulation. I. Detection and masking with narrow-band carriersa). *J. Acoust. Soc. Am.*, 102, 2892–2905. doi: 10.1121/1.420344

- Davis, K. A., & Voigt, H. F. (1997). Evidence of Stimulus-Dependent Correlated Activity in the Dorsal Cochlear Nucleus of Decerebrate Gerbils. *Journal of Neurophysiology*, 78(1), 229–247. doi: 10.1152/jn.1997.78.1.229
- de Boer, E. (1976). On the "residue" and auditory pitch perception. In W. Keidel & W. Neff (Eds.), *Handbook of sensory physiology, vol v-3* (p. 479-583). Berlin: Springer-Verlag.
- de Cheveigné, A. (1993). Separation of concurrent harmonic sounds: Fundamental frequency estimation and a time-domain cancellation model of auditory processing. *J Acoust Soc Am*, 93, 3271-3290. doi: 0.1121/1.405712
- de Cheveigné, A. (1997a). Concurrent vowel identification III: A neural model of harmonic interference cancellation. *J Acoust Soc Am*, 101, 2857-2865. doi: 10.1121/1.419480
- de Cheveigné, A. (1997b). *Ten experiments on vowel segregation* (Tech. Rep.). ATR Human Information Processing Research Labs technical report TR-H-217, https://hal.archives-ouvertes.fr/hal-03090891.
- de Cheveigné, A. (1998). Cancellation model of pitch perception. *J Acoust Soc Am*, 103, 1261-1271. doi: 10.1121/1.423232
- de Cheveigné, A. (1999a). Pitch shifts of mistuned partials: A time-domain model. *J Acoust Soc Am*, 106(2), 887–897. doi:10.1121/1.427104
- de Cheveigné, A. (1999b). Vowel-specific effects in concurrent vowel identification. *J Acoust Soc Am*, *106*, 327-340.
- de Cheveigné, A. (1999c). Waveform interactions and the segregation of concurrent vowels. *J Acoust Soc Am*, 106, 2959-2972. doi: 10.1121/1.428115
- de Cheveigné, A., & Kawahara, H. (1999). Multiple period estimation and pitch perception model. *Speech Communication*, 27, 175-185. doi: 10.1016/S0167-6393(98)00074-0
- de Cheveigné, A., Kawahara, H., Tsuzaki, M., & Aikawa, K. (1997). Concurrent vowel identification i: Effects of relative level and f0 difference. *J Acoust Soc Am*, 101, 2839-2847. doi: 10.1121/1.418517
- de Cheveigné, A., McAdams, S., & Marin, C. (1997). Concurrent vowel identification ii: Effects of phase, harmonicity and task. *J Acoust Soc Am*, 101, 2848-2856. doi: 10.1121/1.419476
- de Cheveigné, A., & Pressnitzer, D. (2006). The case of the missing delay lines: Synthetic delays obtained by cross-channel phase interaction. *J Acoust Soc Am*, 119(6), 3908–11. doi:10.1121/1.2195291
- Deroche, M. L., & Culling, J. F. (2011a). Narrow noise band detection in a complex masker: Masking level difference due to harmonicity. *Hearing Research*, 282(1-2), 225–235. doi:10.1016/j.heares.2011.07.005
- Deroche, M. L., & Culling, J. F. (2011b). Voice segregation by difference in fundamental frequency: Evidence for harmonic cancellation. *J Acoust Soc Am*, 130(5), 2855–2865. doi:10.1121/1.3643812

- Deroche, M. L., & Culling, J. F. (2013). Voice segregation by difference in fundamental frequency: Effect of masker type. *J Acoust Soc Am*, 134(5), EL465–EL470. doi:10.1121/1.4826152
- Deroche, M. L., Culling, J. F., & Chatterjee, M. (2013). Phase effects in masking by harmonic complexes: Speech recognition. *Hearing Research*, *306*, 54–62. doi: 10.1016/j.heares.2013.09.008
- Deroche, M. L., Culling, J. F., & Chatterjee, M. (2014). Phase effects in masking by harmonic complexes: Detection of bands of speech-shaped noise. *J Acoust Soc Am*, 136(5), 2726–2736. doi: 10.1121/1.4896457
- Deroche, M. L., Culling, J. F., Chatterjee, M., & Limb, C. J. (2014a). Roles of the target and masker fundamental frequencies in voice segregation. *J Acoust Soc Am*, 136(3), 1225–1236. doi: 10.1121/1.4890649
- Deroche, M. L., Culling, J. F., Chatterjee, M., & Limb, C. J. (2014b). Speech recognition against harmonic and inharmonic complexes: Spectral dips and periodicity. *J Acoust Soc Am*, *135*(5), 2873–2884. doi:10.1121/1.4870056
- Deroche, M. L., Culling, J. F., Lavandier, M., & Gracco, V. L. (2017). Reverberation limits the release from informational masking obtained in the harmonic and binaural domains. *Attention, Perception, & Psychophysics*, 79(1), 363–379. doi: 10.3758/s13414-016-1207-3
- Divenyi, P. L. (1979). Is pitch a learned attribute of sounds? Two points in support of Terhardt's pitch theory. *J Acoust Soc Am*, 66(4), 1210–1213. doi: 10.1121/1.383317
- Duda, R. O., Hart, P. E., & Stork, D. G. (2012). *Pattern classification*. John Wiley & Sons.
- Duifhuis, H., Willems, L., & Sluyter, R. (1982). Measurement of pitch in speech: an implementation of Goldstein's theory of pitch perception. *J Acoust Soc Am*, 71, 1568-1580. doi: 10.1121/1.387811
- Durlach, N. (1963). Equalization and cancellation theory of binaural masking-level differences. *J. Acoust. Soc. Am.*, *35*, 1206-1218. doi: 10.1121/1.1918675
- Englitz, B., Tolnai, S., Typlt, M., Jost, J., & Rübsamen, R. (2009). Reliability of Synaptic Transmission at the Synapses of Held In Vivo under Acoustic Stimulation. *PLoS ONE*, *4*(10), e7014. doi: 10.1371/journal.pone.0007014
- Ewert, S. D., & Dau, T. (2000). Characterizing frequency selectivity for envelope fluctuations. *J Acoust Soc Am*, 108(3), 1181–1196. doi: 10.1121/1.1288665
- Felix, R. A. I., Gourévitch, B., Gòmez-Àlvarez, M., Leijon, S. C. M., Saldaña, E., & Magnusson, A. K. (2017). Octopus Cells in the Posteroventral Cochlear Nucleus Provide the Main Excitatory Input to the Superior Paraolivary Nucleus. *Frontiers in Neural Circuits*, 11, 37. doi:10.3389/fncir.2017.00037

- Fishman, Y. I., Micheyl, C., & Steinschneider, M. (2013). Neural Representation of Harmonic Complex Tones in Primary Auditory Cortex of the Awake Monkey. *Journal of Neuroscience*, *33*(25), 10312–10323. doi: 10.1523/JNEUROSCI.0020-13.2013
- Fishman, Y. I., & Steinschneider, M. (2010). Neural Correlates of Auditory Scene Analysis Based on Inharmonicity in Monkey Primary Auditory Cortex. *Journal of Neuroscience*, *30*(37), 12480–12494. doi: 10.1523/JNEUROSCI.1780-10.2010
- Fishman, Y. I., Steinschneider, M., & Micheyl, C. (2014). Neural Representation of Concurrent Harmonic Sounds in Monkey Primary Auditory Cortex: Implications for Models of Auditory Scene Analysis. *Journal of Neuroscience*, *34*(37), 12425–12443. doi: 10.1523/JNEUROSCI.0025-14.2014
- Franken, T. P., Bondy, B. J., Haimes, D. B., Goldwyn, J. H., Golding, N. L., Smith, P. H., & Joris, P. X. (2021). Glycinergic axonal inhibition subserves acute spatial sensitivity to sudden increases in sound intensity. *eLife*, *10*, e62183. doi: 10.7554/eLife.62183
- Friston, K. (2018). Does predictive coding have a future? *Nature Neuroscience*, 21(8), 1019–1021. doi: 10.1038/s41593-018-0200-7
- Gábor, D. (1947). Acoustical quanta and the theory of hearing. *Nature*, *159*, 591-594. doi: 10.1038/159591a0
- Gockel, H., Carlyon, R. P., & Plack, C. J. (2009). Further examination of pitch discrimination interference between complex tones containing resolved harmonics. *J Acoust Soc Am*, 125(2), 1059–1066. doi: 10.1121/1.3056568
- Gockel, H., Moore, B. C. J., & Patterson, R. D. (2002). Asymmetry of masking between complex tones and noise: The role of temporal structure and peripheral compression. *J Acoust Soc Am*, 111(6), 2759–2770. doi: 10.1121/1.1480422
- Gockel, H., Moore, B. C. J., & Patterson, R. D. (2003). Asymmetry of masking between complex tones and noise: Partial loudness. *J. Acoust. Soc. Am.*, 114(1), 12. doi: 10.1121/1.1582447
- Gockel, H., Moore, B. C. J., Plack, C. J., & Carlyon, R. P. (2006). Effect of noise on the detectability and fundamental frequency discrimination of complex tones. *J Acoust Soc Am*, *120*(2), 957–965. doi: 10.1121/1.2211408
- Grange, J. A., & Culling, J. F. (2016). The benefit of head orientation to speech intelligibility in noise. *J Acoust Soc Am*, 139(2), 703–712. doi: 10.1121/1.4941655
- Graves, J. E., & Oxenham, A. J. (2019). Pitch discrimination with mixtures of three concurrent harmonic complexes. *J Acoust Soc Am*, 145(4), 2072–2083. doi: 10.1121/1.5096639
- Green, D. M. (1992). On the similarity of two theories of comodulation masking release. *J Acoust Soc Am*, 91(3), 1769–1769. doi: 10.1121/1.402457

- Green, T., & Rosen, S. (2013). Phase effects on the masking of speech by harmonic complexes: Variations with level. *J Acoust Soc Am*, 134(4), 2876–2883. doi: 10.1121/1.4820899
- Grothe, B. (2000). The function of the medial superior olive in small mammals: temporal receptive fields in auditory analysis. *Journal of Comparative Physiology A*, *186*, 413–423. doi: 10.1007/s003590050441
- Guest, D. R., & Oxenham, A. J. (2019). The role of pitch and harmonic cancellation when listening to speech in harmonic background sounds. *J Acoust Soc Am*, 145(5), 3011–3023. doi: 10.1121/1.5102169
- Hafter, E. R., & Saberi, K. (2001). A level of stimulus representation model for auditory detection and attention. *J Acoust Soc Am*, 110(3), 1489–1497. doi: 10.1121/1.1394220
- Hall, J. W., Haggard, M. P., & Fernandes, M. A. (1984). Detection in noise by spectro-temporal pattern analysis. *J Acoust Soc Am*, 76(1), 50–56. doi: 10.1121/1.391005
- Hartmann, W. M., Cariani, P. A., & Colburn, H. S. (2019). Noise edge pitch and models of pitch perception. *J Acoust Soc Am*, *145*(4), 1993–2008. doi: 10.1121/1.5093546
- Hartmann, W. M., & Doty, S. (1996). On the pitches of the components of a complex tone. *J. Acoust. Soc. Am.*, 99, 567-578. doi: 10.1121/1.414514
- Hartmann, W. M., & Goupell, M. J. (2006). Enhancing and unmasking the harmonics of a complex tone. *J. Acoust. Soc. Am.*, 120(4), 2142–2157. doi: 10.1121/1.2228476
- Hartmann, W. M., McAdams, S., & Smith, B. K. (1990). Hearing a mistuned harmonic in an otherwise periodic complex tone. *J. Acoust. Soc. Am.*, 88(4), 1712–1724. doi: 10.1121/1.400246
- Hatfield, G. (2002). Perception as unconscious inference. In D. Heyer & R. Mausfeld (Eds.), *Perception and the physical world: Psychological and philosophical issues in perception* (p. 113-143). John Wiley and Sons, New York.
- Hauth, C. F., & Brand, T. (2018). Modeling Sluggishness in Binaural Unmasking of Speech for Maskers With Time-Varying Interaural Phase Differences. *Trends in Hearing*, 22, 233121651775354. doi: 10.1177/2331216517753547
- Heinz, M. G., Colburn, H. S., & Carney, L. H. (2001). Evaluating Auditory Performance Limits: I. One-Parameter Discrimination Using a Computational Model for the Auditory Nerve. *Neural Computation*, 13(10), 2273–2316. doi: 10.1162/089976601750541804
- Helmholtz, H. (1867). Handbuch der Physiologischen Optik. Leipzig: Voss. (English tranl. 1924 JPC Southall as Treatise on Physiological Optics).
- Holmes, S. D., & Roberts, B. (2012). Pitch shifts on mistuned harmonics in the

- presence and absence of corresponding in-tune components. *J Acoust Soc Am*, 132(3), 1548–1560. doi: 10.1121/1.4740487
- Hu, G., & Wang, D. (2008). Segregation of unvoiced speech from nonspeech interference. *J Acoust Soc Am*, *124*(2), 1306–1319. doi:10.1121/1.2939132
- Huggins, W., & Licklider, J. (1951). Place mechanisms of auditory frequency analysis. *J. Acoust. Soc. Am.*, 23, 290-299. doi: 0.1121/1.1906760
- Imbert, M. (2020). La fin du regard éclairant. Une révolution dans les sciences de la vision au XIe siècle, Ibn al-Haytham. Vrin, Paris.
- Jeffress, L. A. (1948). A place theory of sound localization. *J Comp Physiol Psychol*, 41, 35–39. doi: 10.1037/h0061495
- Jelfs, S., Culling, J. F., & Lavandier, M. (2011). Revision and validation of a binaural model for speech intelligibility in noise. *Hearing Research*, 275(1-2), 96–104. doi: 10.1016/j.heares.2010.12.005
- Jepsen, M. L., Ewert, S. D., & Dau, T. (2008). A computational model of human auditory signal processing and perception. *J. Acoust. Soc. Am.*, 124(1), 422–438. doi: 10.1121/1.2924135
- Joris, P. X., Schreiner, C. E., & Rees, A. (2004). Neural Processing of Amplitude-Modulated Sounds. *Physiological Reviews*, 84(2), 541–577. doi: 10.1152/physrev.00029.2003
- Joris, P. X., & Trussell, L. O. (2018). The Calyx of Held: A Hypothesis on the Need for Reliable Timing in an Intensity-Difference Encoder. *Neuron*, 100, 534–549. doi: 10.1016/j.neuron.2018.10.026
- Joris, P. X., & van der Heijden, M. (2019). Early Binaural Hearing: The Comparison of Temporal Differences at the Two Ears. *Annual Review of Neuroscience*, 42(1), 433–457. doi: 10.1146/annurev-neuro-080317-061925
- Josupeit, A., Schoenmaker, E., Par, S., & Hohmann, V. (2020). Sparse periodicity-based auditory features explain human performance in a spatial multitalker auditory scene analysis task. *European Journal of Neuroscience*, *51*(5), 1353–1363. doi: 10.1111/ejn.13981
- Kadner, A., & Berrebi, A. (2008). Encoding of temporal features of auditory stimuli in the medial nucleus of the trapezoid body and superior paraolivary nucleus of the rat. *Neuroscience*, *151*(3), 868–887. doi: 10.1016/j.neuroscience.2007.11.008
- Kandler, K., Lee, J., & Pecka, M. (2020). The Superior Olivary Complex. In *The Senses: A Comprehensive Reference* (pp. 533–555). Elsevier. doi:10.1016/B978-0-12-805408-6.00021-X
- Kay, R. H., & Matthews, D. R. (1972). On the existence in human auditory pathways of channels selectively tuned to the modulation present in frequency-modulated tones. *The Journal of Physiology*, 225(3), 657–677. doi: 10.1113/jphysiol.1972.sp009962
- Keilson, S. E., Richards, V. M., Wyman, B. T., & Young, E. D. (1997). The

- representation of concurrent vowels in the cat anesthetized ventral cochlear nucleus: Evidence for a periodicity-tagged spectral representation. *J Acoust Soc Am*, 102(2), 1056–1071. doi: 10.1121/1.419859
- Keine, C., Rübsamen, R., & Englitz, B. (2016). Inhibition in the auditory brainstem enhances signal representation and regulates gain in complex acoustic environments. *eLife*, *5*, e19295. doi: 10.7554/eLife.19295
- Kersten, D., Mamassian, P., & Yuille, A. (2004). Object Perception as Bayesian Inference. *Annual Review of Psychology*, 55(1), 271–304. doi: 10.1146/annurev.psych.55.090902.142005
- Klinge, A., Beutelmann, R., & Klump, G. M. (2011). Effect of Harmonicity on the Detection of a Signal in a Complex Masker and on Spatial Release from Masking. *PLoS ONE*, 6(10), e26124. doi: 10.1371/journal.pone.0026124
- Larsen, E., Cedolin, L., & Delgutte, B. (2008). Pitch Representations in the Auditory Nerve: Two Concurrent Complex Tones. *Journal of Neurophysiology*, 100(3), 1301–1319. doi:10.1152/jn.01361.2007
- Laudanski, J., Zheng, Y., & Brette, R. (2014). A Structural Theory of Pitch. *eneuro*, *1*(1), ENEURO.0033–14.2014. doi: 10.1523/ENEURO.0033-14.2014
- Lavandier, M., Jelfs, S., Culling, J. F., Watkins, A. J., Raimond, A. P., & Makin, S. J. (2012). Binaural prediction of speech intelligibility in reverberant rooms with multiple noise sources. *J Acoust Soc Am*, *131*(1), 218–231. doi: 10.1121/1.3662075
- Lea, A. (1992). Auditory Models of Vowel Perception, unpublished doctoral thesis, University of Nottingham.
- Leclère, T., Lavandier, M., & Deroche, M. L. (2017). The intelligibility of speech in a harmonic masker varying in fundamental frequency contour, broadband temporal envelope, and spatial location. *Hearing Research*, *350*, 1–10. doi: 10.1016/j.heares.2017.03.012
- Licklider, J. C. R. (1951). A duplex theory of pitch perception. *Experientia*, 7, 128-134. doi: 10.1007/BF02156143
- Licklider, J. C. R. (1959). Three auditory theories. In S. Koch (Ed.), *Psychology: A study of a science* (pp. 41–144). Mcgraw-Hill.
- Logan, B. F. J. (1977). Information in the zero crossings of bandpass signals. *Bell System Technical Journal*, *56*(4), 487-510. doi: https://doi.org/10.1002/j.1538-7305.1977.tb00522.x
- Lyon, R. (1983, 1988). A computational model of binaural localization and separation. In W. Richards (Ed.), *Natural computation* (p. 319-327). Cambridge, Mass: MIT Press. (reprinted from Proc. ICASSP 83, 1148-1151.)
- Lyon, R. (1984). Computational models of neural auditory processing. In *IEEE ICASSP* (p. 36.1.(1-4)).
- Maxwell, B. N., Richards, V. M., & Carney, L. H. (2020). Neural fluctuation cues

- for simultaneous notched-noise masking and profile-analysis tasks: Insights from model midbrain responses. *J Acoust Soc Am*, *147*(5), 3523–3537. doi: 10.1121/10.0001226
- McDermott, J., & Hauser, M. (2004). Are consonant intervals music to their ears? Spontaneous acoustic preferences in a nonhuman primate. *Cognition*, 94(2), B11–B21. doi: 10.1016/j.cognition.2004.04.004
- McDermott, J. H., Lehr, A. J., & Oxenham, A. J. (2010). Individual Differences Reveal the Basis of Consonance. *Current Biology*, 20(11), 1035–1041. doi: 10.1016/j.cub.2010.04.019
- McDermott, J. H., Schultz, A. F., Undurraga, E. A., & Godoy, R. A. (2016). Indifference to dissonance in native Amazonians reveals cultural variation in music perception. *Nature*, *535*(7613), 547–550. doi: 10.1038/nature18635
- McKeown, D. J. (1992). Perception of concurrent vowels: The effect of varying their relative level. *Speech Communication*, 11(1), 1–13. doi: 10.1016/0167-6393(92)90059-G
- McKeown, D. J., & Patterson, R. D. (1995). The time course of auditory segregation: Concurrent vowels that vary in duration. *J Acoust Soc Am*, 98(4), 1866–1877. doi: 10.1121/1.413373
- Mc Laughlin, M., van der Heijden, M., & Joris, P. X. (2008). How Secure Is In Vivo Synaptic Transmission at the Calyx of Held? *Journal of Neuroscience*, 28(41), 10206–10219. doi: 10.1523/JNEUROSCI.2735-08.2008
- McPherson, M. J., Dolan, S. E., Durango, A., Ossandon, T., Valdès, J., Undurraga, E. A., Jacoby, N., Godoy, R. A., & McDermott, J. H. (2020). Perceptual fusion of musical notes by native Amazonians suggests universal representations of musical intervals. *Nature Communications*, *11*(1), 2786. doi: 10.1038/s41467-020-16448-6
- McPherson, M. J., Grace, R. C., & McDermott, J. H. (2020). Harmonicity aids hearing in noise. *bioRxiv*. doi: 10.1101/2020.05.07.082511
- Meddis, R., & Hewitt, M. (1992). Modeling the identification of concurrent vowels with different fundamental frequencies. *J. Acoust. Soc. Am.*, 91, 233-245. doi: 10.1121/1.402767
- Meddis, R., & Hewitt, M. J. (1991). Virtual pitch and phase sensitivity of a computer model of the auditory periphery. I: Pitch identification. *J Acoust Soc Am*, 89(6), 2866–2882. doi: 10.1121/1.400725
- Meyer, G. F., Plante, F., & Berthommier, F. (1997). Segregation of concurrent speech with the reassigned spectrum. In 1997 IEEE International Conference on Acoustics, Speech, and Signal Processing (Vol. 2, p. 1203-1206 vol.2). doi: 10.1109/ICASSP.1997.596160
- Micheyl, C., Bernstein, J. G. W., & Oxenham, A. J. (2006). Detection and F0 discrimination of harmonic complex tones in the presence of competing tones or noise. *J Acoust Soc Am*, 120(3), 1493–1505. doi: 10.1121/

- 1.2221396
- Micheyl, C., Keebler, M. V., & Oxenham, A. J. (2010). Pitch perception for mixtures of spectrally overlapping harmonic complex tones. *J. Acoust. Soc. Am.*, 128(1), 257–269. doi: 10.1121/1.3372751
- Micheyl, C., & Oxenham, A. J. (2010). Pitch, harmonicity and concurrent sound segregation: Psychoacoustical and neurophysiological findings. *Hearing Research*, 266(1-2), 36–51. doi:10.1016/j.heares.2009.09.012
- Minden, V., Pehlevan, C., & Chklovskii, D. B. (2018). Biologically Plausible Online Principal Component Analysis Without Recurrent Neural Dynamics. *IEEE 52nd Asilomar Conference on Signals Systems and Computers*, 8. doi: 10.1109/ACSSC.2018.8645109
- Moore, B. C. J. (2003). *An introduction to the psychology of hearing*. London: Academic Press.
- Moore, B. C. J., & Glasberg, B. (1983). Suggested formulae for calculating auditory-filter bandwidths and excitation patterns. *J. Acoust. Soc. Am.*, 74, 750-753. doi: 10.1121/1.389861
- Moore, B. C. J., Glasberg, B. R., & Peters, R. W. (1986). Thresholds for hearing mistuned partials as separate tones in harmonic complexes. *J Acoust Soc Am*, 80(2), 479–483. doi: 10.1121/1.394043
- Moore, B. C. J., Glasberg, B. R., Plack, C. J., & Biswas, A. K. (1988). The shape of the ear's temporal window. *J Acoust Soc Am*, 83(3), 1102–1116. doi: 10.1121/1.396055
- Moore, B. C. J., Peters, R. W., & Glasberg, B. R. (1985). Thresholds for the detection of inharmonicity in complex tones. *J Acoust Soc Am*, 77(5), 1861–1867. doi: 10.1121/1.391937
- Needham, K., & Paolini, A. G. (2006). Neural timing, inhibition and the nature of stellate cell interaction in the ventral cochlear nucleus. *Hearing Research*, 216-217, 31–42. doi: 10.1016/j.heares.2006.01.016
- Oh, E. L., & Lutfi, R. A. (2000). Effect of masker harmonicity on informational masking. *J Acoust Soc Am*, 108(2), 706–709. doi: 10.1121/1.429603
- Oja, E. (1982). Simplified neuron model as a principal component analyzer. *Journal of Mathematical Biology*, 15, 267–273. doi:10.1007/BF00275687
- Palmer, A. R. (1990). The representation of the spectra and fundamental frequencies of steady-state single- and double-vowel sounds in the temporal discharge patterns of guinea pig cochlear-nerve fibers. *J Acoust Soc Am*, 88(3), 1412–1426. doi: 10.1121/1.400329
- Parsons, T. (1976). Separation of speech from interfering speech by means of harmonic selection. *J. Acoust. Soc. Am.*, 60(4), 911–918. doi: 0.1121/1.381172
- Piechowiak, T., Ewert, S. D., & Dau, T. (2007). Modeling comodulation masking release using an equalization-cancellation mechanism. *J. Acoust. Soc. Am.*,

- 121(4), 2111–2126. doi: 10.1121/1.2534227
- Plack, C. J., & Moore, B. C. J. (1990). Temporal window shape as a function of frequency and level. *Journal of the Acoustical Society of America*, 87, 2178–2187. doi: 10.1121/1.399185.
- Popham, S., Boebinger, D., Ellis, D. P. W., Kawahara, H., & McDermott, J. H. (2018). Inharmonic speech reveals the role of harmonicity in the cocktail party problem. *Nature Communications*, *9*(1), 2122. doi: 10.1038/s41467 -018-04551-8
- Prud'homme, L., Lavandier, M., & Best, V. (2020). A harmonic-cancellation-based model to predict speech intelligibility against a harmonic masker. *J Acoust Soc Am*, 145(3), 3246–3254. doi:10.1121/1.5101323
- Qiu, J., Wang, H., Lu, J., Zhang, B., & Du, K.-L. (2012). Neural Network Implementations for PCA and its Extensions. *ISRN Artificial Intelligence*, 2012, 1–19. doi: 10.5402/2012/847305
- Roberts, B., & Bregman, A. S. (1991). Effects of the pattern of spectral spacing on the perceptual fusion of harmonics. *J. Acoust. Soc. Am.*, 90(6), 3050–3060. doi: 10.1121/1.401779
- Roberts, B., & Brunstrom, J. M. (1998). Perceptual segregation and pitch shifts of mistuned components in harmonic complexes and in regular inharmonic complexes. *J. Acoust. Soc. Am.*, 104, 2326–2338. doi:10.1121/1.423771
- Roberts, B., & Brunstrom, J. M. (2001). Perceptual fusion and fragmentation of complex tones made inharmonic by applying different degrees of frequency shift and spectral stretch. *J Acoust Soc Am*, 110(5), 2479–2490. doi: 10.1121/1.1410965
- Roberts, B., & Brunstrom, J. M. (2003). Spectral pattern, harmonic relations, and the perceptual grouping of low-numbered components. *J. Acoust. Soc. Am.*, 114(4), 17. doi: 10.1121/1.1605411
- Roberts, B., & Holmes, S. D. (2006). Grouping and the pitch of a mistuned fundamental component: Effects of applying simultaneous multiple mistunings to the other harmonics. *Hearing Research*, 222(1-2), 79–88. doi: 10.1016/j.heares.2006.08.013
- Ruggles, D. R., Freyman, R. L., & Oxenham, A. J. (2014). Influence of Musical Training on Understanding Voiced and Whispered Speech in Noise. *PLOS ONE*, *9*(1), 8. doi: 10.1371/journal.pone
- Saddler, M. R., Gonzalez, R., & McDermott, J. H. (2020). Deep neural network models reveal interplay of peripheral coding and stimulus statistics in pitch perception. *biorRxiv*, 57. doi: 10.1101/2020.11.19.389999
- Sayles, M., Stasiak, A., & Winter, I. M. (2015). Reverberation impairs brainstem temporal representations of voiced vowel sounds: challenging "periodicity-tagged" segregation of competing speech in rooms. *Frontiers in Systems Neuroscience*, 8(248), 19. doi: 10.3389/fnsys.2014.00248

- Scheffers, M. T. M. (1983). *Sifting vowels*. Unpublished doctoral dissertation, Gröningen.
- Scheffers, M. T. M. (1984). Discrimination of fundamental frequency of synthesized vowel sounds in a noise background. *J Acoust Soc Am*, 76(2), 428–434. doi:10.1121/1.391134
- Schmidhuber, J. (2009). Driven by Compression Progress: A Simple Principle Explains Essential Aspects of Subjective Beauty, Novelty, Surprise, Interestingness, Attention, Curiosity, Creativity, Art, Science, Music, Jokes. *arXiv:0812.4360 [cs]*. (arXiv: 0812.4360)
- Schoenmaker, E., Brand, T., & van de Par, S. (2016). The multiple contributions of interaural differences to improved speech intelligibility in multitalker scenarios. *J Acoust Soc Am*, *139*(5), 2589–2603. doi: 10.1121/1.4948568
- Schofield, B. R. (1994). Projections to the cochlear nuclei from principal cells in the medial nucleus of the trapezoid body in guinea pigs. *The Journal of Comparative Neurology*, 344(1), 83–100. doi:10.1002/cne.903440107
- Schroeder, M. (1968). Period histogram and product spectrum: new methods for fundamental-frequency measurement. *J. Acoust. Soc. Am.*, 43, 829-834. doi: 10.1121/1.1910902
- Shackleton, T. M., Meddis, R., & Hewitt, M. J. (1994). The Role of Binaural and Fundamental Frequency Difference cues in the Identification of Concurrently Presented Vowels. *The Quarterly Journal of Experimental Psychology Section A*, 47(3), 545–563. doi: 10.1080/14640749408401127
- Shamma, S. A. (1985). Speech processing in the auditory system II: Lateral inhibition and the central processing of speech evoked activity in the auditory nerve. *J Acoust Soc Am*, 78(5), 1622–1632. doi: 10.1121/1.392800
- Shamma, S. A., & Dutta, K. (2019). Spectro-temporal templates unify the pitch percepts of resolved and unresolved harmonics. *J. Acoust. Soc. Am.*, 15. doi: 10.1121/1.5088504
- Shamma, S. A., Elhilali, M., & Micheyl, C. (2011). Temporal coherence and attention in auditory scene analysis. *Trends in Neurosciences*, *34*(3), 114–123. doi: 10.1016/j.tins.2010.11.002
- Shamma, S. A., & Klein, D. (2000). The case of the missing pitch templates: How harmonic templates emerge in the early auditory system. *J Acoust Soc Am*, 107(5), 2631–2644. doi: 10.1121/1.428649
- Shamma, S. A., & Lorenzi, C. (2013). On the balance of envelope and temporal fine structure in the encoding of speech in the early auditory system. *J Acoust Soc Am*, 133(5), 2818–2833. doi: 10.1121/1.4795783
- Shamma, S. A., Shen, N. M., & Gopalaswamy, P. (1989). Stereausis: binaural processing without neural delays. *J Acoust Soc Am*, 86(3), 989–1006. doi: 10.1121/1.398734
- Shen, Y., & Pearson, D. V. (2019). Efficiency in glimpsing vowel sequences in

- fluctuating makers: Effects of temporal fine structure and temporal regularity. *J Acoust Soc Am*, *145*(4), 2518–2529. doi: 10.1121/1.5098949
- Shera, C. A., Guinan, J. J., & Oxenham, A. J. (2002). Revised estimates of human cochlear tuning from otoacoustic and behavioral measurements. *Proceedings of the National Academy of Sciences*, 99(5), 3318–3323. doi:10.1073/pnas.032675099
- Shore, S. E., Helfert, R. H., Bledsoe, S. C., Altschuler, R. A., & Godfrey, D. A. (1991). Descending projections to the dorsal and ventral divisions of the cochlear nucleus in guinea pig. *Hearing Research*, 52(1), 255–268. doi:10.1016/0378-5955(91)90205-N
- Sinex, D. G., Henderson Sabes, J., & Li, H. (2002). Responses of inferior colliculus neurons to harmonic and mistuned complex tones. *Hearing Research*, *168*(1-2), 150–162. doi: 10.1016/S0378-5955(02)00366-0
- Sinex, D. G., & Li, H. (2007). Responses of Inferior Colliculus Neurons to Double Harmonic Tones. *Journal of Neurophysiology*, 98(6), 3171–3184. doi: 10.1152/jn.00516.2007
- Sinex, D. G., Li, H., & Velenovsky, D. S. (2005). Prevalence of Stereotypical Responses to Mistuned Complex Tones in the Inferior Colliculus. *Journal of Neurophysiology*, *94*(5), 3523–3537. doi: 10.1152/jn.01194.2004
- Slaney, M. (1993). An efficient implementation of the Patterson-Holdsworth auditory filter bank (technical report No. 35). Apple Computer.
- Sorensen, R. (2011). Para-Natural Kinds. In J. K. Campbell, M. O'Rourke, & M. H. Slater (Eds.), *Carving nature at its joints: natural kinds in meta-physics and science* (pp. 113–127). Cambridge, Mass: MIT Press.
- Stasiak, A., Sayles, M., & Winter, I. M. (2018). Perfidious synaptic transmission in the guinea-pig auditory brainstem. *PLOS ONE*, *13*(10), e0203712. doi: 10.1371/journal.pone.0203712
- Stein, A., Ewert, S. D., & Wiegrebe, L. (2005). Perceptual interaction between carrier periodicity and amplitude modulation in broadband stimuli: A comparison of the autocorrelation and modulation-filterbank model. *J Acoust Soc Am*, 118(4), 2470–2481. doi: 10.1121/1.2011427
- Steinmetzger, K., & Rosen, S. (2015). The role of periodicity in perceiving speech in quiet and in background noise. *J Acoust Soc Am*, 138(6), 3586–3599. doi: 10.1121/1.4936945
- Stubbs, R. J., & Summerfield, Q. (1988). Evaluation of two voice-separation algorithms using normal-hearing and hearing-impaired listeners. *J Acoust Soc Am*, 84(4), 1236–1249. doi: 10.1121/1.396624
- Su, Y., & Delgutte, B. (2020). Robust Rate-Place Coding of Resolved Components in Harmonic and Inharmonic Complex Tones in Auditory Midbrain. *The Journal of Neuroscience*, 40(10), 2080–2093. doi: 10.1523/JNEUROSCI.2337-19.2020

- Summerfield, Q., & Assmann, P. F. (1991). Perception of concurrent vowels: Effects of harmonic misalignment and pitch-period asynchrony. *J Acoust Soc Am*, 89(3), 1364–1377. doi: 10.1121/1.400659
- Summerfield, Q., & Culling, J. (1992). Periodicity of maskers not targets determines ease of perceptual segregation using differences in fundamental frequency (abstract). *J Acoust Soc Am*, 92, 2317. doi: 10.1121/1.405031
- Summerfield, Q., Culling, J. F., & Fourcin, A. (1992). Auditory segregation of competing voices: absence of effects of FM or AM coherence. *Philosophical Transactions of the Royal Society of London. Series B: Biological Sciences*, 336(1278), 357–366. doi: 10.1098/rstb.1992.0069
- Summerfield, Q., Foster, J., Gray, S., & Haggard, M. (1981). Perceiving vowels from 'flat spectra'. *J Acoust Soc Am*, 69(S1), S116–S116. doi: 10.1121/1.386490
- Summers, V., & Leek, M. R. (1998). Masking of tones and speech by Schroeder-phase harmonic complexes in normally hearing and hearing-impaired listeners. *Hearing Research*, *118*(1-2), 139–150. doi: 10.1016/S0378-5955(98) 00030-6
- Sumner, C. J., Wells, T. T., Bergevin, C., Sollini, J., Kreft, H. A., Palmer, A. R., Oxenham, A. J., & Shera, C. A. (2018). Mammalian behavior and physiology converge to confirm sharper cochlear tuning in humans. *Proceedings of the National Academy of Sciences*, 115(44), 11322–11326. doi: 10.1073/pnas.1810766115
- Terhardt, E. (1974). Pitch, consonance, and harmony. *J Acoust Soc Am*, 55(5), 1061–1069. doi: 10.1121/1.1914648
- Tollin, D. J., & Yin, T. C. T. (2005). Interaural Phase and Level Difference Sensitivity in Low-Frequency Neurons in the Lateral Superior Olive. *The Journal of Neuroscience*, *25*, 10648–10657. doi: 10.1523/JNEUROSCI.1609-05.2005
- Treurniet, W. C., & Boucher, D. R. (2001). A masking level difference due to harmonicity. *J Acoust Soc Am*, 109(1), 306–320. doi:10.1121/1.1328791
- Verschooten, E., Desloovere, C., & Joris, P. X. (2018). High-resolution frequency tuning but not temporal coding in the human cochlea. *PLOS Biology*, *16*(10), e2005164. doi: 10.1371/journal.pbio.2005164
- Verschooten, E., Shamma, S., Oxenham, A. J., Moore, B. C., Joris, P. X., Heinz, M. G., & Plack, C. J. (2019). The upper frequency limit for the use of phase locking to code temporal fine structure in humans: A compilation of viewpoints. *Hearing Research*, 377, 109–121. doi: 10.1016/j.heares.2019.03.011
- Viemeister, N. F. (1979). Temporal modulation transfer functions based upon modulation thresholds. *J Acoust Soc Am*, 66(5), 1364–1380. doi: 10.1121/1.383531

- Viemeister, N. F., & Wakefield, G. H. (1991). Temporal integration and multiple looks. *J Acoust Soc Am*, 90(2), 858–865. doi: 10.1121/1.401953
- Walker, K. M., Gonzalez, R., Kang, J. Z., McDermott, J. H., & King, A. J. (2019). Across-species differences in pitch perception are consistent with differences in cochlear filtering. *eLife*, 8, e41626. doi: 10.7554/eLife.41626
- Wan, R., Durlach, N. I., & Colburn, H. S. (2014). Application of a short-time version of the Equalization-Cancellation model to speech intelligibility experiments with speech maskers. *J Acoust Soc Am*, 136(2), 768–776. doi: 10.1121/1.4884767
- Wang, D. (2008). Time-Frequency Masking for Speech Separation and Its Potential for Hearing Aid Design. *Trends in Amplification*, 12(4), 332–353. doi: 10.1177/1084713808326455
- Wang, D.-L., & Brown, G. (2006). Computational auditory scene analysis: Principles, algorithms and applications. IEEE Press / Wiley.
- Weintraub, M. (1985). A theory and computational model of auditory monaural sound separation. Unpublished doctoral dissertation, Stanford.
- Whiteford, K. L., Kreft, H. A., & Oxenham, A. J. (2020). The role of cochlear place coding in the perception of frequency modulation. *eLife*, 9, e58468. doi: 10.7554/eLife.58468
- Xie, R., & Manis, P. B. (2013). Target-Specific IPSC Kinetics Promote Temporal Processing in Auditory Parallel Pathways. *Journal of Neuroscience*, *33*(4), 1598–1614. doi: 10.1523/JNEUROSCI.2541-12.2013
- Zheng, Y., & Escabí, M. A. (2013). Proportional spike-timing precision and firing reliability underlie efficient temporal processing of periodicity and envelope shape cues. *Journal of Neurophysiology*, *110*(3), 587–606. doi: 10.1152/jn.01080.2010
- Zwicker, U. (1984). Auditory recognition of diotic and dichotic vowel pairs. Speech Communication, 3(4), 265–277. doi: 10.1016/0167-6393(84) 90023-2