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The fate of the Benue-Congo velar nasal in Bantoid

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Abstract:

Proto-Benue-Congo has been reconstructed both with a simple velar nasal * η and with the velar nasal as part of a * ηg structure. The loss of the simple velar nasal has been noticed as a feature of Bantu languages but has not been studied in other Bantoid groups. This paper considers the lexemes for which this phoneme has been reconstructed and then examines the subsequent development of the velar nasal in several groups. The results vary from preservation in Ekoid to near absence in Bantu. The phonological loss of velar nasals is not surprising, as parallels demonstrate. Rather the question arises of why certain languages only preserve remnants of the original velar nasal, and a partial answer is connected to suffixation and cluster formation.

Keywords: Bantoid, Bantu, velar nasal, reconstruction

1 Introduction

¹ Cf. Hyman (2003: 49), Meeussen (1967: 83, 85), and Meinhof et al. (1932: 32–3).

² Unless otherwise stated, PBC reconstructions are from de Wolf (1971: 53–59) with class markers in parentheses, while PB reconstructions are taken from *Bantu*

Mukarovsky (1976–77: 248) also noted the unusual loss of his Proto-Western Nigritic *n in C $_2$ position in certain PB reconstructions. Then Dimmendaal (1978: 233–34) was the first to state that there is a regular correspondence between stem-medial and stem-final * η in Proto-Upper Cross (PUC) and \emptyset in Proto-Bantu for which he gave several examples.

But besides the Bantu reflexes, what happened to the PBC velar nasal in other Bantoid groups? After highlighting the general problem of velar nasals in Proto-Benue-Congo and Niger Congo, the plan of this article is first to identify stems which can be reconstructed with simple $*\eta$ at the level of Proto-Cross-Congo (PCC) or earlier, and then look in detail at a range of Bantoid language groups for evidence of preservation or loss. Finally, we will look at the Bantu evidence for greater clarity on that subgroup.

The Bantoid data in this paper is cited from publications whose authors vary in their conventions for transcribing or analyzing nasal + velar structures (ηg or ng, prenasalized or not, compound or not). But I will refer to all those structures as complex ηg in contrast to simple η (without any additional component). Likewise, the term " η loss" only refers to a language's loss of the inherited PBC simple phoneme * η not to its loss in a complex structure. It is also important to keep in mind that the presence of simple η in a language today does not mean that the language failed to undergo " η loss", since after the

Lexical Reconstructions 3 (BLR3) (Bastin, Coupez, Mumba & Schadeberg 2002), sometimes with examples from Guthrie (1967–71). The convention of BLR3 (following Meeussen's usual style) is to write *ng for the PB prenasalized velar, but in this paper I use *ng, as had been the custom of Meinhof and Guthrie. Regardless of the convention of the authors cited, I write reconstructed roots without preceding or following hyphens. Letters following PB reconstructions indicate the Guthrie zones where reflexes are attested; the symbol "+" indicates additional zones, that is to say "ABDE+" is a shorthand for zones A, B, D, E and some further letters.

- 3 A description of possible Bantoid branches can be found in Blench (2015), but it must be stated that there is a lack of clear characteristics, much less confirmed innovations, that would distinguish the members of a Bantoid group or many subgroups, including Bantu itself. Although Dakoid is included in Bantoid in Blench (2009), I think much of its core vocabulary has roots divergent from the conventional Bantoid languages.
- 4 To label the ancestor of the Cross River and Bantoid languages, I use the term Proto-Cross-Congo (rather than Proto-Bantoid-Cross) to maintain consistency with the pattern Niger-Congo and Benue-Congo and to avoid confusion with PBC, the abbreviation for Proto-Benue-Congo. PCC reconstructions are mine.

loss of inherited * η the simple velar nasal often arose anew from the reduction of * ηg or other consonants.

2 The velar nasal in early Benue-Congo or Niger-Congo

There is not yet a list of reconstructions for PCC or Proto-Bantoid, so we must begin at an earlier stage. What Benue-Congo roots might be candidates for studying the reflexes of PBC simple $*\eta$ in Bantoid? De Wolf (1971) was focused on PBC noun classes rather than phonology and he provides no description of how he reconstructed his tentative PBC phonemic system, which includes both $*\eta$ and $*\eta g$, or even how he reconstructed roots. Nevertheless, de Wolf's main list of tentative PBC noun roots includes twenty roots with simple $*\eta$ and nine roots with $*\eta g$. Only some of these have Bantoid reflexes. In his list of PBC phonemes, Blench (2004) reconstructs *m, n, n, and *N (presumably an unspecified nasal), but he also uses $*\eta$ in his Eastern Benue-Congo reconstructions (e.g. 253 $\#ko\eta$ 'hill') and $*\eta g$ in the Bantoid-Cross ones (e.g. 329 $\#ka\eta ga$ 'guinea-fowl'). Sometimes these match PB reconstructions, sometimes they do not.

In fact, in Niger-Congo in general, the complex * ηg is not very common (mainly Bantu and the Atlantic or Mel languages). Among his 600+ pseudo-reconstructions of Proto-Western-Nigritic (covering most of the groupings of current Niger-Congo), Mukarovsky (1976–77)

⁵ De Wolf (1971: 51–59), grouped by gender. Those with ηg are *kwenge 'cheek', * ηg wana 'child', * $ka\eta ga$ 'guinea fowl', * $ba\eta ga$ 'jaw', * $kpa\eta ga$ 'mat', * $pe\eta ga$ 'shoulder', * $kwo\eta ga$ 'spear', * $ta\eta ga$ 'stick, whip', * $ga\eta go$ 'tooth'. In addition, there are occasional incidental reconstructions throughout the volume.

includes eight with a simple velar nasal or ηw complex (all initial only). The PWN complex *ng appears 26 times (only in C_2 position), often as an alternative, and often without examples for $ng/\eta g$ outside Bantu. Even the presence of ηg in a sample language is not very indicative. For example, one of the few languages that Mukarovsky regularly cites with ηg is Gonja, a North Guang language of the Kwa group, but Snider (1989) uses η or $\eta \eta$, e.g. e- $pe\eta i$ 'sun' instead of Mukarovsky's \dot{e} - $pe\eta gi$. In any case, Snider (1990: 10) derives North Guang * $\eta \eta$ from Proto-Guang * ηk , so Guang will not provide any confirmation for reconstructing Proto-Niger-Congo * ηg . Yet, for the widely attested stem of 'guinea fowl' (PWN *kuang, PBC * $ka\eta ga$, PB * $ka\eta ga$), other than the Bantu and Benue-Congo reconstructions, Mukarovsky's only reflexes with * ηg are a Gonja form and possibly Kisi (Mel) $ka\eta gb$ -u 'kind of dove'.

The paradox is that PB has * ηg but no simple * η , but beyond Bantu there is plenty of simple η but little evidence for * ηg . There is a similar deficit of evidence in languages outside Bantu for the other nasal complexes (*mb and *nd) reconstructed by Mukarovsky and de Wolf, but there are at least numerous oppositions in Bantu of *m/*b/*mb

⁶ Mukarovsky (1976–77) discusses η at 1.116–17, and the "denasalization of * ηm " in some languages at 1.142–46 (later he uses * ηw in listing those same roots). Mukarovsky writes his PWN pseudo-constructions with capital letters, but for ease I have written them with miniscules. Those with * η : * ηiu (* ηium , * ηu -) 'drink', * ηun (* ηunk) 'smell', * ηwa 'scratch', * ηwa (* ηwya , * ηwu na) 'nose', * ηwa (*) 'smoke', * ηwa (*) 'smoke', * ηwa (*) 'smohe', * ηwa (*) 'bee; honey'.

⁷ PWN roots with *ng: *baga (*banga) 'jaw', *bung 'waterhole, river', *beng 'dislike, hate', *bhang 'shine brightly', *bhung 'roar', *cang 'charm', *gwàng 'hang up', *jing 'build', *kuang 'guinea-fowl', *kung 'bend (eel)', *kwúng 'gather up', *liang 'hang down, dangle', *lingi 'water', *mún (mung) 'suck', *mang (*mak) 'astonish', *ní (*níang) 'elephant tusk', *pin (*ping) 'lie across', *pok (*pong) 'knead', *pek (*peng) 'be bent sideways', *ping 'plait', *phán (*pháng) 'cutlass', *phuk (*phung) 'blow, breathe', *tagi (*tangi) 'kind of trap', *tang (*tiang) 'be first, in front', *ting 'cut (off)', *tung 'basket', *tsuang? (*tuang) 'stick'.

⁸ De Wolf has only a few reconstructions with *mb or *nd and, as usual, the evidence for them is not clear. For example, none of his examples for *kondi 'moon, month' or *tende 'palm tree' have /nd/, so presumably they are based on PB *gòndè and *téndé respectively. For *kumba 'pig' (no PB cognate), he cites the Upper Cross River forms Mbembe ò-kómbà and Humono è-kómbá/ì-. But /mb/ is very rare inside a morpheme in those languages, and the Okom variant akku 'pig' suggests that the Mbembe forms okkómba, akkómba involve a clitic. Dimmendaal does not reconstruct *mb, *nd or *ng for PUC.

and *n/*d/*nd. Without PB reconstructions, which heavily influenced Mukarovsky and de Wolf, it is quite possible that PWN and PBC might have been reconstructed with just * η , and not * ηg . Two general solutions to the puzzle are obvious: 1) a widespread shift of * $\eta g > \eta$ except in Bantu, 9 or 2) only early * η (without * ηg) and then Bantu innovated * $\eta > \eta g$. From the phylogenetic viewpoint, it seems unusual for so many branches of Volta-Congo to lose a phonemic feature independently. But the fact that the other PB nasal complexes look solid makes an early * ηg plausible typologically.

How solid is the Bantu evidence for non-initial * ηg ? The various NC₂ structures are regularly attested in hundreds of languages from all zones, regardless of tones or surrounding vowels. There is an occasional tendency toward nasal complexes inside the history of the Bantu group, e.g. early Bantu *yàb-uk- 'cross river' > Eastern Bantu *yàmb-uk. But there is really nothing internal to justify the case for a uniform change like PBC * η > PB * ηg (and in certain cases PB * \emptyset). 10 Bantu languages differ in this regard from the neighboring Grassfields languages, where NC, structures are frequent but no NC, structures have been reconstructed for Proto-Grassfields. 11 In general, the simplification of consonants in C2 and the limitation of coda consonants are widespread phenomena in the Nigeria-Cameroon area and are often connected with the loss of V₂ but do not require it. Some Bantu languages in the area were also affected by these changes, for example, PB *ng is fairly consistently preserved by Sawabantu languages like Duala (A24), as well as by Tuki (Mbam A601), but not usually by Tunen (A44) or the Manenguba group: e.g. PB *gàngá 'root' > Duala mwangá and Tuki ongangá but Tunen mòkana and Akossi (A15c) $\dot{\eta}$ -k $\dot{\alpha}\eta$. The change * $\eta g > \eta$ (and sometimes further η $> \emptyset$) is also a feature of many B70–80 languages.

⁹ If both existed earlier, this would have been a merger of * ηg and η . For the change * $\eta g > \eta$ and general background on the velar nasal in Germanic languages, see Bailey (2019).

¹⁰ Among the C_2 consonants for a hypothetical Proto-Potou-Akanic-Bantu (PPAB), Stewart (2002: 209) lists mb, nt, ηg , ηk , and n (but not m or η). He states, "the simple nasal n is presumed to go back diachronically to a prenasalized stop nd" (so the occasional PB *nd is a non-systematic innovation when it occurs, as is Proto-Akanic * $\eta < n$. But his * ηg is early, e.g. PPAB * $ti\eta gi$ 'vein' > PB * $ti\eta gi$ - $tilde{li}$.

¹¹ The apparent exception PGr *sàm´bà(l)´ 'seven' (PB càmbàdì, càmbòàdì BLR 8433) is a compound formation.

In summary, there is reason to wonder about the early history of pre-nasalized consonants or nasal complexes at the stages before Proto-Bantoid. But putting aside these speculations about the phonemes of Proto-Niger-Congo and Proto-Benue-Congo, this paper will continue with the standard assumption that PBC had both * ηg and * η and that PB * ηg is a reliable indicator of the former. The question under study then is to what extent PBC had simple * η and what its fate was in Bantoid languages.

2.1 PBC or PCC reconstructions with simple velar nasals

If they were in fact distinct, PBC * ηg and * η merged in early Cross River in favor of * η , e.g. PB * $k\acute{a}\eta g$ 'fry, roast' ~ PUC * $k\acute{a}\eta\grave{a}$, PLC * $k\acute{a}\eta$. So, by themselves the forty PUC roots with * η in Dimmendaal (1978) do not distinguish any difference in source. But he devised a simple test for distinguishing them by listing six cognate roots for which PUC * η has a corresponding * \emptyset in Guthrie's PB, thus demonstrating the phenomenon of simple η loss in Bantu: 'knee', 'egg', 'ear', 'louse', 'root/hair', 'faeces'. The Bantu comparisons are needed because Cross River * η is ambiguous in itself. Likewise, reconstruction based on Bantoid alone is complicated by languages where contemporary η can have a variety of sources. Accordingly, the evidence of PB is critical for distinguishing PCC * η and * ηg .

For this analysis, I have further added several roots from other sources that might be relevant. Proto-Lower Cross (PLC), Proto-Cross River and Proto-Ogoni reconstructions are from Connell (1991, 2025a), Proto-Western-Nigritic (PWN) reconstructions from Mukarovsky (1976–1977), Benue-Congo Reconstructions (BCR) from Blench (2004), Proto-Jukunoid (PJ) from Shimizu (1980), Proto-Plateau (PP) from Gerhardt (1983: 139–154), Proto-Grassfields (PGr) from Hyman (1979). Also cited when available is data from the Benue-Congo Comparative Wordlist (BCCW) collected in Williamson & Shimizu (1968) and Williamson (1973). Many contemporary language forms are those collected by the Grassfields Bantu Working Group (GBWG) in the 1970s or in the Bantu database accompanying Grollemund et al. (2015).

We can create a list of possible PBC or PCC forms with a simple velar nasal by filtering the various lists above for roots where a PB

¹² Dimmendaal 1978: 233–34. For the velar nasal in specific Upper Cross languages, with reconstructions of PUC, see Sterk (n.d.: 71–2).

form lacks the velar nasal found in the PBC, PUC, or PLC reconstruction. Simple velar nasals have not been reconstructed at the beginning of PBC roots. ¹³ I have separated the root-medial (1) and root-final examples (2), because assessing final velar nasals is often more complicated. Some roots do not have good Bantoid evidence for a final consonant, and in some forms it is possible that the reconstructed final velar nasal is based on an original suffixed class marker (especially likely to be preserved on monosyllables). Since all PB nouns are reconstructed with a vowel coda, it is also possible that the lack of final velar nasals is due to a general loss of all PB final consonants.

(1) PBC or PCC proto-forms with velar nasal in medial position

'ear' PBC *tuŋi (ku/a)

PB *tớì 15/6, 5/6 (BLR 3030)

The velar nasal is well attested in Cross River (PUC * $tt\acute{o}\eta(i)$, PLC * \acute{u} - $tz\acute{\eta}$ /a-) and somewhat in Jukunoid, Plateau and Western Gur (BCCW 31, PJ * $t\acute{o}\eta$ (ku/a), PWN 556 $th\acute{u}i$, $th\acute{u}$). We will see that Bantoid languages sometimes show what look like frozen *-lr or *-kv suffixes, both frequently seen as class markers in Gur forms.

'ashes' PBC *toŋ (de Wolf 1971: 71) PB *tó, *túé 3, 14 (BLR 2954)

Despite de Wolf, Bantu forms support a CVCV structure for PCC. The velar nasal is well attested in Plateau, Jukunoid and Cross River (BCCW 5, PJ *tóŋ (u-/i-), PWN 560 *thún). Across Benue-Congo and Bantoid, the most common vowel in this word is $\mathfrak I$ (so PUC *tíŋ, PLC *ń-tɔŋ́). Guthrie's reconstruction of *túé (C.S.1810.5, 1769), followed by BLR, is based on forms like Songe (L23) *e-twe*, which helpfully provide V_2 , but the glide in those forms only suggests an unspecified rounded back vowel and all the examples with vowels in zones ABD have o or $\mathfrak I$.

'hair, root, vein' PUC *dɪŋa (dò-/ɪ)

PB *dì 3/4, *dia 3 'root, fibre' (BLR 1003)

The PUC reconstruction is supported by PLC *5-l $\delta\eta$ 'root', but not PWN 344 *lful, ful 'root', PJ *dzin(u/i) 'hair' or Proto-Central-Jukunoid * $d\acute{e}n$ 'root', PLC * $d\acute{e}t$ 'hair'. Unfortunately, it is not easy to be sure about Bantoid cognates because the root has

¹³ Even a prenasalized velar appears initially in only one of de Wolf's stems: PBC *ŋgwana/*bana 'child', where the nasal in the singular is clearly a remnant of the class prefix *mu-.

metaphorical semantics, which seem to overlap with possible by-forms or other roots:

- 1. PUC *tına (dò-/ı) 'root, hair', PWN 515 *tin 'root (trunk)', PB tínà 'root, base of tree trunk banana plant'.
- 2. PBC *duŋu (ú/ti) 'cord / rope', *digi (ú/i) 'cord / rope / liana'; PJ *diT, *diK (u/i) 'rope'; PUC -dyíkì, PLC *ó-lók; PB *dígì 3/4 'string' in zones CDE+, PB *dí 11 'string' BGH+. The two PBC proto-forms for 'cord, rope' look suspiciously similar especially since the velar nasal in *duŋu is probably just a velar, as in Cross River.
- 3. PB *cìŋgà 'string; hair (on body)'.
- 4. Also, for 'root', Bantoid languages often use the cognates of PB *gàngá.

'knee' PUC *dúŋí (dè-/dà-) PB *dúí 5/6 (BLR 1239)

The velar nasal is likewise reconstructed for PLC * $\acute{\epsilon}$ - $l\acute{5}\eta$ /n- and BCR 220 # $r\acute{u}\eta\grave{u}$ (with forms from Plateau and Tarokoid). PWN 363 * $l\acute{u}nku$ - (* $d\acute{u}nku$ -). There is limited data for PWN 109 *dui. The basis of De Wolf's reconstruction of PBC *duno (*li/*a) is unclear since none of the forms he cites have C₂ or V₂. Perhaps he was making a connection with the Eastern Bantu * $d\grave{v}\eta g\grave{o}$ 'knee' EGJL+ (BLR 1217) but BLR derives that from the verb 'join'. PBC *kudu (li/a) may reflect this root with a * $k\grave{v}$ class prefix for body parts.

(2) PBC or PCC proto-forms with velar nasal **in final position** 'dew' PBC *min (ma)
PB *mè 3, 5, 11 (BLR 2158)

The velar nasal is found in Plateau reconstructions but the word was not included in the usual Cross River lists for reconstruction. Blench (2022) gives common Plateau #myene, with n as a Plateau innovation. PWN 366 main, mina. A possible conflation exists with stems for 'water': PBC ni (main); or PLC main, PGr main.

'egg' PBC *kiŋ, *tiŋ PB *gí 5/6 (BLR 1368)

Reconstruction of the velar nasal for PBC is uncertain at best. Cf. PJ *kyì (ri/a), PP*tii and *zeŋ, PWN 122 *gíla, BCCW #33. From PUC (Dimmendaal 1978: 297) *kkèŋí, PUC (Sterk n.d.:

80) *`-kkèddí and PLC *ý-kíɛn, Connell reconstructs Proto-Cross-River *kiɛn. These forms and the Bantoid forms discussed below often appear to have the relic of a class 5 marker.

'excrement' PBC *biŋ (a)

PB *bîî 13, 6 (BLR 6425)

The velar nasal is seen in reconstructions of Plateau and Jukunoid (PJ *byíŋ /a-), but for Proto-Cross-River, Connell reconstructs *bín (cf. PUC (Dimmendaal 1978: 256) *`-biŋ, PUC (Sterk n.d.: 72) *`-bín, PLC possibly *fíŋ or *fíD). PWN 32 *bin. BCR 98 #mi. Protoforms for Mambiloid and Grassfields are reconstructed with final *d, see below.

'louse' PUC *dáŋ (è-/i-), PLC *ó-láŋ / i-PB *dá 9/10 (BLR 780)

Dimmendaal (1978: 67) and Sterk (n.d.: 34–5) discuss the problem that * η is absent in this stem in a number of Central Upper Cross languages. But Northern Jos has * $la\eta$ (bi/i). BCR 240 #daNi is based on Tarokoid (Sur) dani, the PLC reconstruction and Nizaa (Mambiloid) $l\bar{a}m$. No entries for 'louse' in PWN or BCCW, and de Wolf only offers PBC *i-kodu/*i-. I have not found other Bantoid cognates.

'mosquito' PUC *boŋ, PLC *é-bɔŋ PB *bύ 9/10, 11/10 (BLR 279)

Also Proto-Plateau 2 (Jaba) i-buŋ. Sterk's reconstruction of PUC * $^{\cdot}$ -bbɔʻŋŋ \mathring{V} suggests a suffix is at issue. The velar nasal is not supported by PWN 53 *búne, BCR233 #boN (in fact, none of the three forms cited there has any nasal).

'wing' PBC *pabaŋ, *babaŋ (li/a) PB *pàpá 5/6, 11/10, *bàbá (BLR 2407)

The velar nasal is seen in some Plateau languages but is not reconstructed for Plateau groups. It is, however, reconstructed for PUC * $b\acute{a}b\grave{a}p$ (Dimmendaal 1978: 258) and perhaps with a suffix in PUC * $\dot{b}a\acute{b}a\acute{p}$ (Sterk cited by Connell 2025a) but not PLC * \acute{u} - $b\^{a}$ / m-. See also BCCW 101; PWN 419 *papa 'shoulder, wing'. This word, and related verbs to 'flap wings', may have an onomatopoetic dimension that complicates reconstruction.

Altogether, these ten roots above provide a few good cases for reconstructing a PBC or PCC simple velar nasal, although only four are medial, and in most of the word-final cases the evidence is rather

weak. It should be noted that all of these are nouns, partly because de Wolf's list only includes nouns but also because the great majority of PUC reconstructions with velar nasals are nouns.

2.2 Unlikely or uncertain cases of PBC *ŋ and *ŋg

Few BC languages have pre-nasalized or compound NC structures, so de Wolf apparently reconstructed $*\eta g$ on the basis of the Bantu data like that in Table 1.

Table 1. Proto-Benue-Congo forms in *ŋg

	PBC	PB
'guineafowl'	*kaŋga (ì/î)	*káŋgà 9/10 (BLR 1720)
ʻjaw'	*baŋga (ku/î)	*báŋgá 11/10 (BLR 108)
'spear'	*kwoŋga (li/a)	*gòŋgá 5/6 (BLR 1448)

Thus, we should be able to pair PBC * ηg with PB * ηg and PBC * η with PB * ϕ . But it is well known that many of de Wolf's reconstructions deserve reconsideration. Based on PB or other Bantoid reflexes, it seems that some PBC forms, listed in (3), should actually have different types of velars than those reconstructed by de Wolf, or none at all.

(3) Disputable PBC or PCC proto-forms

'shoulder' PBC *peŋga (ku/a)

PB *bègà 5/6 (pègà 5/6) (BLR 139)

De Wolf (1971: 63, 158) only gives two uncertain reflexes and PBC * ηg looks unlikely here in general: PWN 41 *buaka (bueka); PP2 (Jaba) gu-pek / a-, PGr * $mb\`ek$ (-a?); Ekoid: Balep $\grave{\varepsilon}$ - $p\varepsilon g$, Ejagham $ef\^a\eta$, Ekajuk $\acute{\varepsilon}$ - $b\acute{\sigma} g$.

'tooth' PBC *gango (\pm) (li/a)

PB **gègò* 5/6 'molar tooth' (BLR 1356)

De Wolf (1971: 62) only gives one possible example Central Kambari áá-ngá/á. Cf. PWN 200 *kéka.

'buffalo, bush cow' PBC *pon (ì/î)

PB *bògó 9/10 (BLR 258)

De Wolf gives several examples with final -m (so also PUC * $p\acute{o}m$ 'cow') but none with a velar nasal. The Bantoid evidence suggests either * ηg or possibly *g: PGr * $mb\acute{o}\eta$ '; in Yemne-Kimbi, Bu $mb\acute{o}l\bar{a}/b\grave{m}b\acute{o}l\bar{a}$ (1/2) and Koshin $mb\acute{o}\eta/b\grave{o}mb\acute{o}\eta$ (1/2); the

Ekoid-Mbe evidence is discussed later. If PBC * ηg , perhaps the PB form in classes 9/10 reflects some dissimilation of double NC structures – to which the varying C_1 of this difficult word may also contribute. Cf. BCCW 46.

'blood' PBC *luŋ (ma)

PB *dúŋgό 6 (BLR 4488)

De Wolf's examples do not show a velar nasal (or ηg), but we will see below that the Bantoid data supports PBC * ηg . There are other 'blood' roots that might be relevant for some of the Bantoid data: 1) PBC *zi (ma), PGr *cl', PB *gida (5) 6; 2) PJ * $yi\eta$ (ma), PUC * $yi\eta$, PB * $ji\eta ga$ 6, but perhaps some forms are related to the previous series; 3) PGr * $l\acute{e}m$ ` looks like a cognate of PUC * $d\grave{e}$ but is more likely from PBC * $lu\eta$ with the class 6 marker *-ma suffixed.

'charcoal' PBC *kalaŋ (li/a) PB *kádà 5/6 (BLR 1662)

There is no strong evidence for * η in Plateau or Jukunoid and Connell (2025a) reconstructs Proto-Cross River * $k\acute{a}DV$. In any case, a putative C₃ in this form would be at risk in most languages. Cf. BCCW #16, PGr * $k\acute{e}$ `. There is also the problem of the PB pair * $k\acute{a}da\eta g$ /* $k\acute{a}\eta g$ 'roast'.

'leopard' PBC *kpoŋi (ì/î)

Without a clear PB descendant, it is hard to know whether to reconstruct PBC * η or * ηg , although the latter seems more likely. De Wolf's velar nasal seems to be based on a form in Kainji (Anaguta \dot{u} - $kp\dot{e}\eta/\dot{i}$), and three Bendi forms: Alege \dot{i} - $kp\dot{o}\eta$, Boki \dot{i} - $kpa\eta/\dot{i}$ - and Bete \dot{u} - $kp\dot{o}\eta/\dot{i}$, which are ambiguous. PBC * $kpo\eta$ i has some resemblance to PB * $g\dot{o}$ i 9 'leopard' (BLR 7154), but there must have been several terms for wild cats, and there is no velar nasal in PJ * $k\dot{u}$ (?)(i/i), PUC * $kp\dot{e}$ and PLC * \acute{e} - $kp\dot{e}$ /*i-, PWN 150 * $gw\dot{i}$ l. Cf. BCCW 56. Some groups like Ekoid have both roots (or stems of the same root?).

'mouth, lip' PBC *nuŋa (ú/ti)

PB *nòà 12, *nyòà (BLR 4709)

De Wolf only cites Kagoro (Plateau) \dot{u} - $n\acute{u}\eta$ with the velar nasal, and there is limited support for a velar nasal from Cross River. PLC * \acute{t} - $n\acute{u}\grave{a}$ argues against the reconstruction of a velar nasal, as does PUC * $m\grave{a}$ (often forms in $mm\grave{a}$), which is likely from *nua

with a labialization of the nasal by the rounded vowel. Proto-Ogoni *ngegā is ambiguous at best. See also PWN 404 níuma, núa. In general, this root has been seen as a likely denominative of 'drink' (cf. PB *nyó (BLR 7047), PWN 406 *niu(m), *nu), and *-a or a class affix seems a more likely suffix than *-na. But note occasional second nasals in Bantu forms for 'drink' like Nugunu (A622) nyóɛna, Mangisa (A63) a-nyuŋ, Eton (A71) mè-núŋ.

'throat/voice' PBC *toŋo (ì/î)

The velar nasal in Plateau 2, Ekoid, Bendi, and PLC * \dot{u} - $t\acute{o}\eta$ 'neck' is ambiguous. The noun is apparently not known in Bantu but the Mambiloid and Grassfields evidence discussed below suggests the root was *tongo. In that case, perhaps related is the verb seen in PB * $t\acute{o}\eta g$ 'crow, groan' and Nchane (Beboid) $th\bar{o}\eta\bar{e}$ 'crow (v)' (Boutwell 2020). Williamson (1989: 250–54) considers # $to\eta$ a Benue-Congo innovation.

'sheep' PBC *tiane (*bu/*i)

De Wolf's evidence is very weak, including Tiv *ì-yɔ̂ngò*. PUC *tama and PJ *tam better fit the fragmentary Bantu data: *tààmà 9 (BLR 9217) cited from zone J and forms in -tomba found in various A70 languages. Bendi data discussed below.

'tooth' PBC *sana (li/a)

There are still several other roots with a simple velar nasal in de Wolf, but they lack clear PB correspondences.¹⁴ Because of the complexi-

¹⁴ Not discussed in this article are: PBC *taŋ (* \dot{u} /*ba) 'thief', *kpaŋe (*ku/*a) 'canoe/boat', *ziŋin (* \dot{i} /* \dot{i}) 'guineafowl', *toŋi (* \dot{l} i) 'honey', *kiaŋ (*ku/*a) 'thing', *kodog (*u/*i) 'throat / voice', *mbag (*u/*i) 'rain, rainy weather', *tiegu (*u/*ba) 'witch' and *tiegu (*u) 'witchcraft'.

ties of numbers as a separate topic, I have also not treated 'four' in this article. However, the velar nasal reconstructed for Cross River (PUC *nànì; *nàn, PLC *iniàn) is absent not only in Bantoid languages (including PB *nài) but also in Jukunoid, Kainji, and most Platoid languages (Pozdniakov 2018).

3 The velar nasal in various Bantoid groups

To summarize so far, based on data outside Bantoid, we have about ten lexemes for which we might want to reconstruct a simple velar nasal for the proto-Bantoid stage. That is not a lot, but enough to examine the Bantoid evidence, with an emphasis on the four stems with medial $*\eta$ ('ashes', 'ear', 'hair/root' and 'knee'). As we go along, we can re-assess some of the problematic reconstructions.

3.1 Ekoid-Mbe

We begin with the Ekoid-Mbe group which provides the fullest data. These languages regularly have a consonantal reflex of PBC * ηg , but they show a difference in reflexes of the PBC simple * η . Ekoid varieties like Ejagham and Efutop regularly preserve the simple velar nasal, but Mbe does not, as shown in Table 2. Included are Proto-Ekoid-Mbe (PEM) reconstructions from John Watters (p.c.).¹⁵

		-		
Language	'ashes'	'ear'	'excrement'	'knee'
PBC	*toŋ	*tuŋi	*biŋ	*dúŋí
PEM	*tôŋ	*tôŋ	*bîŋ	*rúŋ~lúŋ (?)
Ejagham	a-tôŋ	o-tûŋ	à-b î ŋ	a-rúŋ, ê-r ú ŋ
Efutop	ǹ-tûŋ	ò-tâàŋ	à-bîŋ	è-dâòŋ
Mbe	bé-tùè (pl.)	ó-tūÈ	a-bí	lé-lù

Table 2. Ekoid-Mbe evidence for preservation of a velar nasal

The word for 'root' (PUC *dma - PB *di 3/4, *dia 3) is one more example where Ekoid has a velar nasal (N. Etung i-yinj) but Mbe does not (e-li/be-1). An alternative stem is seen in

¹⁵ Unless otherwise stated, Ekoid data comes from Blench (2014), which is based on Ejagham data from John Watters and David Crabb's comparative wordlist. The Mbe data comes from the unpublished wordlists collected by Otronyi et al. (2009) or Blench (2014).

¹⁶ For *d > Etung y, Mbe l, cf. PB * $d\acute{o}m$ 'bite' \sim Etung yûm, Mbe $lw\acute{o}m$.

PEM * $r\check{u}g$ (?) 'root' (Ejagham $ir\grave{u}k$, Efutop \grave{n} - $du\check{s}g/\grave{a}$ -) which might suggest a velar plosive rather than velar nasal in PBC * $du\eta u$ 'cord/rope', as discussed above.

Table 3. Ekoid-Mbe evi	dence for lacl	k of inherited	velar nasal

Language	'egg'	'wing'	'mouth'	'mosquito, bee'
PBC	*kiŋ,	*pabaŋ,	*пила	[PUC *boŋ,
	*tiŋ	*babaŋ		PLC *é-bɔŋ]
PEM	*kĭ	(*)- $paparepsilon$	nyò	búm
Ejagham	eji	efaβe	ὴμὸ/ὸμὸ (GBWG)	т́-bэ́т
Efutop	è-zí	_	_	m̀-bú
Mbe	lèkī:	épūpūrù	<i>ú</i> ɲūŋ	_

Since Ejagham and Efutop seem reliable indicators of PBC velar nasals, it is noteworthy that velar nasals are missing in Ekoid reflexes of some of de Wolf's PBC reconstructions, as seen in Table 3. No Ekoid languages have a final nasal in the roots for 'egg' or 'wing'. For 'mouth', Ekoid evidence suggests there is no * η in PBC, but Mbe has a surprising velar nasal. For 'mosquito, bee', the final m in Ejagham and the missing final nasal in Efutop suggest an incorporated class 9/10 marker rather than a lost velar nasal.

It must be emphasized that the lack of velar nasals in the Mbe forms above is not because Mbe does not have that phoneme. Rather, Mbe is like Bantu in losing the simple velar nasal but showing a consonantal reflex of PBC * ηg , e.g. PB * $k\acute{a}\eta g$ 'fry, roast' ~ Efutop $ka\acute{a}\eta\grave{a}$, Mbe $y\acute{a}\eta$; cf. PB * $b\acute{u}mb$ 'mould pottery' ~ Efutop $m\grave{o}\eta$, Mbe $mw\acute{o}\eta$. Accordingly, the presence of a velar nasal in both the Ekoid and Mbe forms of a word, together with PB * ηg , is a good indicator that the relevant PCC reconstruction should be * ηg . This distinction suggests three revisions to PBC reconstructions (Table 4).

Language	Phonemes	'blood'	'horn, tusk'	'cow, buffalo'
PBC	various	*luŋ	*tano	*роŋ
PCC	*ŋg (vs. *ŋ)			
Ejagham	ŋ	a-gy ú ŋ	ń-táŋ	m-fóŋ
Efutop	ŋ	à-lớŋ	ǹ-tə́ŋá	ὴ-fáŋ
Mbe	ŋ (vs. ø)	bē-lūŋ	ņ-táŋ	ṇ-pūɔ̄ŋ (lìbˈɛ̀)
PB	*ŋg (vs. *ø)	*dúŋgó	*tóŋgʊ	[*bògó]

Table 4. Series of PCC *ng rather than simple velar nasal

At the PCC stage, it certainly seems that all three of these roots should be reconstructed with $*\eta g$, and accordingly at the PBC stage too, rather than de Wolf's proposals.

Likewise, the final consonant in Mbe \hat{n} -júkp \hat{n} 'leopard' suggests * ηg in PBC * $kpo\eta i$ 'leopard'. But the reconstruction of 'leopard' is uncertain (as discussed above), especially since most Ekoid varieties have cognates of PB * $gonalta \hat{n}$ 9 'leopard', e.g. Ejagham \hat{m} -gbe/ \hat{o} or Ekparabong $\hat{\eta}$ - $k\hat{o}i/b\hat{o}$ -.

Overall, it seems that Proto-Ekoid-Mbe inherited a distinction between simple * η and complex * ηg . Ekoid later merged them, while Mbe lost the velar nasal. So Ekoid and Mbe forms help us establish distinct reconstructions and confirm medial * η in 'ashes', 'ear', 'hair/root' and 'knee', as well as support a final consonant in 'excrement' (perhaps earlier *d, cf. PGr * $b\dot{l}d$).

3.2 Bendi

Closest to Ekoid geographically, and perhaps linguistically, are the Bendi languages. Like most Bantoid groups, Bendi languages have simple velar nasals as $\rm C_2$ reflexes of PBC * $\eta g \sim \rm PB$ * ηg (Table 5). ¹⁷

Table 5. Correspondences of PB * ηg with Bendi η

	•	
Meaning	PB	Bendi data
'guineafowl'	*káŋgà	kě-kâŋ (Bokyi), kù-kâŋ (Bisu)
'tooth'	*còŋgà	-∫áŋ (Bokyi), う-∫áŋ (Alege)
ʻblood'	*dúŋgó	$\bar{\varepsilon}$ - $r\bar{u}\eta$ (Ukwortung), èb $\hat{\varepsilon}$ - $n\bar{u}\eta$ (Bekwara)
'horn'	*tóŋgʊ	í-t ^ị úŋ (Ukwortung)

¹⁷ Bendi group examples are from Blench (2010) and the wordlists in Otronyi et al. (n.d.). The ηg in C_2 of Bendi \dot{u} - ηg $\dot{\eta} g$ $\dot{i}\dot{e}$ 'woman' (where other varieties have g, $\eta \eta$, or η) is merely due to reduplication.

But a number of Bendi varieties have final m where η is expected, e.g. Bekwara itf^wom 'horn'. On the other hand, the final η in Ukwortung $\hat{u}l^i\hat{u}\eta$ 'dry season' (PGr * $l\hat{u}m$ `, Bekwara $\hat{u}j\hat{\sigma}m$) and $\hat{t}^i\hat{\epsilon}\eta$ 'heart' (cf. PB $tim\hat{a}$, Bekwara $rit^i\hat{\epsilon}m$) is surprising. Although Ukwortung has a number of words with final m (e.g. $ik\hat{u}m$ 'corpse'), it can also elide it altogether, e.g. inal 'animal' (cf. PB nnam, Bekwara inal). Until these coda rules or developments are clarified, it is useful to cite multiple Bendi varieties in cases of divergence. For example, we find differing nasals in Bendi as well as in Benue-Congo reconstructions for 'sheep': Ukwortung $inal} inal$ (cf. PBC * $inal} inal} inal}$

In 'ear' and 'ashes', the velar nasal is seen in almost all varieties (4).

- (4) Bendi simple velar nasals as reflexes of PBC velar nasals:
 - a. 'ear' (PBC *tuŋi (ku/a) PB *tới 15/6) Bete ko-tʃoŋ /a-, cf. Bisu ko-ton/a-
 - b. 'ash(es)' (PBC *toŋ PB *túé 3)Bete à-tyúŋ, Bisu à-tʃûŋ

Other Bendi words which look like inheritances of simple velar nasals are 'knee' (e.g. Mbube $r\bar{\iota}$ - $r\acute{u}\eta/\bar{e}$ - $r\acute{u}\eta$) and 'root' (e.g. Okworogung $\bar{\iota}$ - $d\hat{a}\eta$), presumably cognate with PUC * $dr\eta a$ (PB * $d\hat{\iota}$ 3/4, *dia 3) where Ekoid also has a velar nasal but Mbe does not.

Bete \hat{u} - $kp\hat{a}\eta/\hat{i}$ - 'leopard', ki- $tu\eta$ 'neck' and Bokyi $mpo\eta$ 'dwarf cow' are ambiguous for * ηg or * η but are most likely reflexes of * ηg , as discussed above. There is no C₂ in Bekwara $\hat{o}\eta\hat{u}$ 'mouth', or Bete $\hat{l}\hat{i}$ - $\hat{t}\hat{l}$ '(egg', presumably from PBC * $ti\eta$ (li/a). Bokyi bu- $byibaba\eta$ 'wing' is exceptional for the final velar nasal (cf. Bete $k\hat{u}$ - $b\hat{i}b\hat{a}$), but perhaps the extra syllables are an onomatopoetic effect of flapping wings. Cf. the reduplication in Ekoid varieties like Balep $\hat{\epsilon}$ - $kp\hat{o}\eta mkp\hat{o}\eta/b\hat{\epsilon}$ -.

In general, the Bendi group behaves like the neighboring Ekoid, preserving a velar nasal in the key words for 'ear', 'ashes', 'knee', and omitting it in 'mouth'. There are no signs of suffixation.

3.3 Tivoid group and its neighbors

Blench (2016) has rightly raised questions about the membership of the Tivoid group. I have even more doubts than he does, since I do not think the presence of both prefixes and suffixes in these languages is sufficient for grouping. I would limit the name Tivoid to

the two groups he calls Central Tivoid A and B, each of which can be further divided. Near them, Esimbi and Ugarə (Mesaka) look like distinct other languages. Of those so-called Tivoid languages west of the Mambila Plateau, based on lexical isoglosses and more frequent relics of nasal prefixes, I would distinguish the Batu group (Blench's Northern Tivoid), and two other languages Njwande (Bitare) and Buru, which are perhaps related to or influenced by Batu.¹⁸

3.3.3 Tivoid

Generally, Central Tivoid languages have CV(C) roots with a varying range of prefixes and suffixes. First, we should observe some forms that look like inherited * ηg , e.g. Caka-Batanga oganga 'root' (PB * $g\grave{a}ng\acute{a}$), $iyang\jmath$ 'spear' (PB * $g\grave{o}ng\acute{a}$), $ating\varepsilon$ 'count (v)' (PB * $t\acute{a}ng$) and Tiv aungwa 'hear' (PB * $j\acute{t}(\eta)gua$). But some caution is warranted because most Central Tivoid A verbs rarely have ηg in any position, and noun forms suggest that the NC combinations in Tivoid can sometimes be expanded versions of a simple nasal C_2 , either by phonological processes or the inclusion of suffixes, for example we seem to see * $d \sim *nd$ (PB $b\acute{e}\grave{e}d\grave{e}$ 'breast', Caka-Batanga ibanda), as well as * $m \sim mb$ (PBC *lemi 'tongue' > Otanga $olemba\jmath$, Tiv $n\acute{o}mb\acute{o}r$). ¹⁹

Whether current ηg in Tivoid is a preservation or a secondary development, the reflexes of the inherited * η in 'ear' and 'ashes' in (5) and (6) mostly show loss.

(5) Tivoid examples for 'ear' (PBC *tuŋi (ku/a) – PB *tới 15/6) Central Tivoid B: Caka-Batanga ɔtu, Eman ɔto Central Tivoid A north: Tiv utox, Iyive kátɔ́k/átɔ́, Otanga oto Central Tivoid A other: Evand kʷɔ́tɔ́ŋgɔ́k/átɔ̄, Oliti otuŋ, Baceve ɔtuŋo

Tivoid languages often preserve both class prefixes and old suffixes, hence the Iyive form historically analyzable as $k\delta$ - $t\delta$ -k. The final consonant in Tiv utox probably also historically reflects the Tiv class marker ku as it does in itiox 'head' (PB * $t\dot{v}$ e). The unusual example of a singular-plural contrast in Evand $k^w \delta t \delta \eta g \delta k / \delta t \bar{\delta}$ looks like a reflex

 $^{18\,}$ Data on the languages grouped with Tivoid are from Blench (2016) and Koops & Blench (2010).

¹⁹ This phenomenon may also be seen in varieties of Mambiloid (Connell 2025b), e.g. 'navel': Mvur Mambila kubil, but Cambap Mambila $k\acute{o}mb\bar{o}n$, Oumyari Wawa $t f\acute{o}mb\bar{a}i$ (PB * $k\acute{o}b\acute{u}$ 5/6, Tiv i- $comb\acute{o}$).

of *kv- $tv\eta$ -kv/*a- $tv\eta$ -a 15/6 (with an additional *-k added once the original one assimilated to g). Apparently, the class 15 suffix -k in the singular provided a consonant cluster environment that preserved the nasal C_2 , but the vocalic plural marker did not. The Oliti and Baceve forms look like reduced forms of the Evand form.

(6) Tivoid examples for 'ash(es)' (PBC *toŋ – PB *túé 3) Oliti mutumu, Tiv ituɛm, Iyive mutwim

No variety preserves a velar nasal. The Oliti form looks rather like what we would expect from a class 3 noun with a *mu- prefix and a related suffix (or perhaps a *wu concord). Evand mɔtumɛlam is similar with an additional suffix. Independent diphthongization is rare in these languages, so Tiv ituɛm seems to faithfully preserve the same vowels as PB *túé, and Iyive mutwim is similar but with glide formation.

Tivoid provides no good evidence for the preservation of a velar nasal in other candidates like 'mouth', 'excrement', and 'dew'. For 'egg', almost all languages have prefixed forms of gi, ge, or ki, with apparent remnants of class 5 suffixes seen in Baceve $\varepsilon dzin\varepsilon$, Oliti $\varepsilon dzi\eta$.

3.3.2 Other languages sometimes labelled Tivoid

Esimbi seems to maintain *ηg sometimes, e.g. ɔ-ghəŋgə 'root' and mɛ-yuŋgu 'blood', but note also təŋə 'count' (PB *táŋg). If Esimbi atáŋgə 'moon' is related (as Indo-European words for 'month' are from *meh₁ 'measure'), then there might be a difference of intervocalic and final reflexes (or suffixes?). In contrast, the inherited simple velar is lost in Esimbi, e.g. mɔɔtù 'ashes', ó-to/ɔ́-to 'ear'. In short, Esimbi patterns with some of the Central Tivoid languages.

Likewise, Ugare (Mesaka) distinguishes inherited * $\eta g > \eta$ (ú-tǎŋ 'moon', u-gaŋ 'root') from inherited * η (ú[†]tô 'ear', vatu 'ash').

The Batu group is not always consistent but has fewer NC complexes, e.g. Afi $\eta g \acute{o} g \grave{o} n / \grave{a} k \acute{o} \eta$ 'spear' and $mb \acute{a} n \bar{o} n / \acute{a} b \acute{a} \bar{n}$ 'breast', but these reflexes contrast with the absence of consonants in the words for 'ear': Afi $nd \acute{u} n / \acute{a} n \acute{d} \acute{o}$, Njwande out / atu. Afi $m \bar{u} t \bar{a} m w \bar{u}$ 'ashes' has either a *- $m \upsilon$ or *- $w \upsilon$ suffix, which makes it unclear whether a C₂ consonant is preserved. There is a final -n in Kamino and Afi \not{o} - $n \acute{u} n / \acute{a}$ - 'knee' but lacking in some other varieties, so it might be a frozen class marker, like the suffix seen in 'ear'. Kamino $g \hat{u}$ 'leopard' shows no C₂. Likewise in Buru, we can distinguish the reflexes of * ηg (e.g.

 \bar{e} - $g\acute{a}\eta$ 'root', e- $\theta\bar{a}\eta$ 'horn') from the loss of simple * η (e.g. \bar{e} - $t\acute{e}$ 'ear', e- $n\acute{u}/a$ -nu 'knee').

In summary, if the distinctions seen in Esimbi, Ugare, Buru and some Tivoid languages are systematic, then they might also be used to distinguish PCC * ηg from * η , in the way that PB and Mbe can be. The only Tivoid+ relics of the simple PCC * η seem to be in the word for 'ear' in a few languages where the velar nasal was protected by a CV suffix.

3.4 Mambiloid

Mambiloid deserves a separate section because Endresen (1990/1991: 191) noted three nouns where "The coda phoneme η in Pre-Nizaa B corresponds to Proto-Bantu \emptyset . Once again a Proto-Bantu innovation: loss of a phoneme that is retained in Nizaa"; see (7).

- (7) Endresen's correspondence of Pre-Nizaa-B * η to PB * φ^{20}
 - a. 'urine': Nizaa tſúŋ¯ PB cờ
 - b. 'ear': Nizaa twấã, Pre-Nizaa B *tÓηa PB *tớì
 - c. 'mouth': Nizaa $n\tilde{u}\tilde{u}$, Pre-Nizaa * $n\tilde{u}\eta$ - \hat{U} PB * $n\tilde{v}a$.

²⁰ I have cited modern Nizaa forms based on the phonetic transcriptions in Endresen (1990/1991) or Endresen (1992). The orthography of modern Nizaa indicates the nasalization of long nasalized vowels by the use of a subsequent η , e.g. $tw\hat{a}\eta$ [twã:] 'ear', which is to be distinguished from the actual velar nasal following short vowels, e.g. $t\hat{y}\hat{u}\eta$ 'urine'.

²¹ A reviewer points out that the cognacy of Nizaa $t \hat{y} \hat{u} \hat{y} \sim PB c \hat{v}$ is made suspect by the unexplained difference in tone. For the Proto-Bantu reconstruction, the forms with *b (*c\hat{u}b\hat{a}\hat{u}\text{ 'urine'} BH and *c\hat{u}b(ad) 'urinate' BDJFHM) look as early as those without (*c\hat{v}\text{ DFJKLMR}, with Eastern D).

We can add 'ashes' (PBC *toŋ – PB *túé) to the list of words with velar nasal preservations in Mambiloid based on Ndoro $\bar{a}t\bar{o}\eta\bar{a}$, Oumyari (Wawa) $t\bar{o}\bar{o}\eta g\bar{o}$, Vute $t\bar{u}u\eta$ and Mbaw $at\hat{o}\eta$. But other than the exception of 'ashes' and the Nizaa forms above, Mambiloid languages seem to have lost the inherited velar nasal, which is not surprising since even in Nizaa the evidence is mostly the trace of a nasalized vowel. Rather, we find Ndoro $\eta w\bar{u}$ 'mouth' and Oumyari $nuk/num\dot{o}$, and even 'ear' is reported without the nasal in Ndoro $t\bar{o}$, Oumyari $t\bar{o}/t\dot{o}-m\bar{o}$, and Vute $t\dot{q}$. This loss is in contrast to the frequent preservation of the pre-nasalized velar PBC *tg0 ~ PB *tg1, as seen in PB *tag1 'fry' ~ Cambap (Mambila) tag2, Ndung (Kwanja) tag3, Oumyari tag3, oumyari tag3, or PB *tag4, or PB *tag6, as seen in PB *tag6, or PB *tag6, or PB *tag7, oumyari tag8, oumyari tag9, 'gather' ~ Vute tag9.

Mambiloid does not seem to provide any evidence supporting the PBC or PCC reconstruction of a simple velar nasal in final position. Rather, for PBC *miŋ 'dew', we find Ndoro ā-mɛ 'dew'; for PBC *biŋ 'excrement', Ndoro bí (Proto-Mambiloid *bid, Connell (p.c.)); for PUC *bɔŋ 'mosquito', Len mòk; and for PBC *kalaŋ 'charcoal', Ndoro tʃānā, Maberem tʃanı and Vute kāŋkāàr.

In the case of de Wolf's reconstruction of PBC *toŋo (\hat{i}/\hat{t}) 'throat, voice', we do not have a PB reflex to distinguish whether the root has a velar nasal or a prenasalized velar. In Mambiloid, however, forms like Ndoro $t\bar{o}\eta g\bar{o}$, Maberem $t\dot{o}\eta g\dot{o}$ 'neck', and Mbaw $t\dot{o}k$ 'throat, neck' indicate that the PBC and Proto-Mambiloid forms contained a prenasalized velar stop or consonant cluster. This would allow us to connect this root with the verb PB *tóŋg 'crow, groan', Nizaa $t\dot{o}\ddot{o}$ 'crow (v)', Vute $t\dot{o}\eta$ -.

So, Mambiloid languages apparently preserve a few traces of the PBC velar nasal. Some varieties may also provide helpful evidence to distinguish the inherited velar nasal from nasals of other origins.

3.5 Beboid and Yemne-Kimbi

In examining some other Bantoid branches, we find that once again the words for 'ashes' and 'ear' offer solid examples of preserved simple velar nasals, but few other words do. On the whole, it becomes

²² Unless otherwise specified, in this section Ndoro forms are cited from Connell & Blench (2014), Mbaw forms from GBWG, Wawa-Oumyari from Martin (2012), Vute from Thwing (1987) and other Mambiloid varieties from Connell (2025b).

apparent that suffixed class markers were an important factor for the preservation of the inherited simple velar nasal.²³

In Beboid and Yemne-Kimbi, the complex PBC * ηg (reflected in PB * ηg) has reflexes in simple η , as seen in (8).

(8) Beboid and Yemne-Kimbi velar nasals corresponding to PB *ŋg
PB: *gàŋgá 'root', PB *dúŋgó 'blood', *tóŋgu 'horn'
Pabaid Sari gwán 'root', Mwngang khlóán 'blood',

Beboid: Sari gwéŋ 'root', Mungong kàlúúŋ 'blood'

Noni ke-tóŋ 'horn'

Yemne-Kimbi: Ajumbu əgoŋ 'root'

The words 'ear' and 'ashes' in (9) and (10) also maintain the inherited simple velar nasal, but apparently as part of old clusters:

(9) 'ear' (PBC *tuŋi (ku/a) – PB *tớì 15/6) Beboid: Saari kintɔ́ɔ̄η, Cung ntônē

Yemne-Kimbi: Abar kətənə, Bu kətunwə

The Yemne-Kimbi and Beboid examples show various prefixes and suffixes, or suffix influence. Noni (Beboid) has a helpful doublet: in addition to the pair ke- $t\acute{u}/bi$ - $t\acute{e}m$, it has a reduplicated byform $k\grave{e}$ - $ntu\eta$ - $t\acute{u}\eta/b\grave{i}$ -, which seems to preserve the velar nasal. Likewise, Koshin $k\bar{a}$ - $t\acute{u}$ has a diminutive $f\bar{a}$ - $t\acute{u}n$ -f, which is likely derived from * $f\bar{a}$ - $t\acute{u}\eta$ -f.

(10) 'ash(es)' (PBC *toŋ – PB *túé 3)

Beboid: Sari $t\bar{a}\eta\bar{\varepsilon}$, Mungong $t\bar{a}\eta$

Yemne-Kimbi: Bu dʒəŋ

As regular as these velar nasals may look, there are a number of surprises that suggest the frequent presence of suffixes or suffixal remnants. Beboid languages usually lose V_2 ; however, we see Sari $t\bar{a}\eta\bar{\epsilon}$ and Akweto $t\bar{a}\eta g\bar{\epsilon}$ (GBWG) with V_2 and a surprising ηg that look like developments from some suffixal cluster. Noni (Beboid) distinguishes final m (ηam 'animal'), n (tin 'five'), and η (tin 'horn'), so Noni taan 'ashes' is also unexpected unless a suffix influenced it.

Elsewhere, for 'knee': Sari nunu/nuun 5/6, Bukwe $\acute{n}ny\~u$ (Beboid) and Kenyang (Nyang) $n\acute{\epsilon}-n\acute{\epsilon}n/m\acute{a}$. For 'excrement', Mekaf (Beboid) $m-b\grave{\partial}-m$ historically shows the same double affixes for cl. 6a that Noni

²³ Unless otherwise indicated, Beboid is from the Eastern Beboid lists in Blench (n.d.) or GBWG, Esimbi and Ugare from Blench (2016), Yemne-Kimbi from the Western Beboid lists in Blench (n.d.) but I follow Good et al. (2011) for the term Yemne-Kimbi and name Ajumbu (instead of Mbu').

has: prefix m- and suffix -m. The alveolar nasals in Yemne-Kimbi cognates like Koshin $t\bar{o}mbin$ (13) and Missong $k\bar{i}ban/b\bar{i}ban$ (7/8) are more likely reflexes of the *d seen in PGr *bid and Mundabli mbyil (Voll 2017). Only Ajumbu mwain 'dew' might reflect PBC *min, with labialization from the class 3 prefix.

I have found no useful evidence in Beboid for velar nasal preservation in other words. In Noni, final η (from * ηg) is preserved in class 6 but dropped in the singular form, e.g. $l\bar{e}e/\bar{e}l\bar{\partial}\eta$ 'pumpkin' (PB $densurement{e}\eta g$) and $g5\bar{o}/\epsilon g5\eta$ 'spear' (PB *gonga) (Hombert 1980: 90). From this pattern, Mekaf has generalized the pattern \emptyset /- η for a dozen nouns in classes 5/6, e.g. $de/da\eta$ 'pumpkin' and $t\bar{u}/t\bar{u}\eta$ 'horn', and so it is not possible to deduce any original value from Mekaf $gi/ga\eta$ 'egg' or $\eta u/\eta u\eta$ 'knee'. See also the discussion of 'egg' under Bantu below. There is also a final nasal or nasalization on the vowel in 'mosquito' in some Beboid varieties, e.g. Mbuk bwen ~ bwen 5 'mosquito' and $\beta u/\beta u$ 'blood-sucking fly' 9/10 (Tschonghongei 2018).

In sum, unlike Ekoid and Bendi, which have several words preserving the PCC simple velar nasal, the other Bantoid branches we have discussed (Tivoid and neighbors, Mambiloid, Beboid and Yemne-Kimbi) provide solid examples only in 'ear' and 'ashes', often in forms with suffixal influence. Otherwise the velar nasal in these languages reflects PCC *ŋg or some other consonant structure. As we continue the Bantoid migration southeastward, we will see that even that short list diminishes, with Grassfields showing an inherited velar nasal only in suffixed forms of 'ear'. Even more reduced, Tikar has nasalized vowel reflexes of PBC *ŋg, e.g. ngán 'guinea fowl' and yán 'roast', but apparently no remnant of *ŋ in dwí 'knee', ndî 'root', and mè myì 'dew'; the ancestor of the glottal stop in ywí 'ear' and mè byi' 'excrement' is unclear (Jackson 1988).

3.6 Grassfields

As the branch closest to Bantu, the Grassfields data is particularly relevant for assessing the stage of the loss of velar nasals. Final velar nasals are very common in Grassfields languages today where they

²⁴ I do not include Dakoid in Bantoid, but for comparison relevant Dong forms from Blench (2009) for this study would be *tuŋwa* 'ashes' (the suffix *-wa* might have an origin as a class 3 concord, cf. Daka *tóòmáa*), *utuŋ* 'ear', *ruŋ* 'knee' (cf. Gaa *àlúŋa*), *vīī* 'excrement' (nasalization apparently not due to loss of final *m*, cf. *wɔm* 'husband').

arose as reductions of various consonants including original prenasalized velars, which are preserved in Bantu. See the examples in (11).

(11) Grassfields simple velar nasals as correspondences of PB *ŋg

PB *gòŋgá 'spear' – PGr *yòŋ`

PB *gàngá 'root' – PGr *yàn´

PB *dàngí 'bamboo' – PGr *dìní

PB *bòngó 'brain' – PGr *bón

PB *táng 'count' - PGr *tán

But there is very little evidence of the survival of the inherited velar nasal in Grassfields. The only good example is 'ear' (12) which has an unusual PGr structure with suffix -li.²⁵

(12) 'ear' (PBC *tuŋi (ku/a) – PB *tới 5/6, 15/6 – PGr *túŋ-li)

East Gr: Bandjoun tăŋ, Bafou làtùŋǵ, Baba táŋlá, Bazou tòŋú,

Mankon àtôŋná

Momo: Njen àtòŋá, Moghamo-Batibo àtòŋí

Ring: Kom àtúŋló, Oku kētóōlé, Aghem kótúŋō, Isu kótúŋī²⁶

SW Gr: Tanka etoŋi, Menka ətəŋnə, Atong etəŋni, Busam ıtoŋni

Several things are of note here. First, these Grassfields forms are typically in genders 5/6 or 7/8. Secondly, V_2 of noun roots is typically not preserved in Grassfields, so the final vowel here is almost surely from the suffixation of the class 5 concord * d_I -, as Hyman and Elias reconstructed. Thirdly, there is apocope of the final vowel of the root, creating the rare clusters ηl and ηn – which were then often simplified to simple η . It looks like the velar nasal survived here only as part of a consonant cluster, because in other Grassfields words an inherited velar nasal is uncertain at best.

We have seen that most Bantoid branches have some velar nasal reflexes in the word for 'ashes': PBC *toŋ, PB *túé 3, PGr *tóe (?). The evidence is limited for Grassfields (hence Hyman's question mark)

²⁵ Grassfields forms (including Otang) are cited from the Grassfields Bantu Working Group (GBWG), and Babanki forms from Akumbu (2008). For the name and forms of "Southwest Grassfields", see Blench (2010a), although key lexical differences make me wonder if these languages should actually be included under the Grassfields umbrella.

²⁶ Babanki *kàtyítyí?/àtyítyí?* (7/8) is clearly reduplicated but does it show loss of velar nasal or is the final glottal stop a remnant?

²⁷ For other relics of a suffixed concord in Ring languages, see Akumbu and Wills (2024).

because other roots are used for 'ashes' in most branches except in Southwest Grassfields, e.g. Obang *mátúá* (6a), and some Momo, e.g. Lower Mundani *mímót*. The fricative in Atong (SW) *ntuyɔ* is a common Western Grassfields hiatus filler, resolving the hiatus seen in the closely related Menka *ontfúó*. In the case of preservation, we would expect the velar nasal itself, as in Atong *etɔ́ŋní* 'ear'. The important point is that nowhere in Grassfields is a velar nasal observed for this word, only the hiatus (or a hiatus filler) from the loss of the velar nasal. In contrast to forms of 'ashes' in other Bantoid branches, there is no evidence in Grassfields of suffixation, and also no evidence of preservation.

To reinforce the point, it is worth a look at other lexemes which show velar nasals in some Bantoid languages, but not in Grassfields. First are some examples with reconstructed medial velar nasals (13).

(13) Words without medial velar nasal reflexes in Grassfields 'knee' PBC *duno ~ PCC *dúní – PB *dúí 5/6 – PGr *lúl No velar nasals observed in Grassfields, rather forms that share the Bantu loss, e.g. Babanki àlwí.

'mouth' PBC *nuna – PB *nòà

Grassfields branches regularly use another root (PGr *cùl`), but SW Grassfields does have reflexes of the PBC root, e.g. Bantakpa $\varepsilon \mu u$. For Ambele, Jungraithmayr et al. (1975) give $\dot{\varepsilon}$ - $n\dot{o}(\eta)/\dot{a}$ -but his Ambele words for 'ear' and 'ash' just have -tO with no C₂, while Blench lists Ambele enu 'mouth'.

Likewise, in (14) Grassfields provides several examples of words without the velar nasals reconstructed by de Wolf in final position:

- (14) Words without final velar nasal reflexes in Grassfields 'charcoal' PBC *kalaŋ (li/a) PB *kádà 5/6 PGr *ké` No velar nasal found.
 - 'dew' PBC *miŋ (ma) PB *mè 3, 5 PGr *mÙk, *mèk No velar nasal is observed in Grassfields examples. Rather Hyman reconstructed a final stop based on the frequent C_2 seen in forms like Fe'efe'e (Bamileke) mù?°, Adere-Dzodinka (Nkambe) $m \grave{a}k$ °, Oku (Ring) $\bar{\imath} \jmath w \hat{a}k$, Aghem (Ring) $t m \acute{a} ? \grave{a}$, Bum (Ring) $\bar{a} t \bar{a} m \acute{u} A k \grave{a}$. These words for 'dew' look like forms of 'water' (PGr *mò') with a diminutive suffix *ka.

'egg' PBC * $ki\eta$, * $ti\eta$ (li/a) – PB *gi 5/6

Eastern Grassfields, Momo and Ring use forms of *bum 'egg', with the exception of the curious Ngwaw/Ngwo (Momo) àkôn/èkôn (7/8). But Ngwaw àtìmé/ètìmé (7/8) 'ear' suggests some nasals in that language are of uncertain history.

'excrements' PBC *biŋ (a-) – PB *bi \hat{i} 13 – PGr *b $\hat{i}d$

No velar nasal is found in reflexes of this root, rather final stops: Moghamo (Momo) *tíbít*, Baleng (Bamileke) *ndzét*, Bamoun (Nun) *mbít* (note the stop also in Bamoun *tít* 'ear').

'wing' PBC *pabaŋ, *babaŋ (li/a) – PB *pàpá 5/6, *bàbá PGr *bàb`-li

No velar nasal is observed in plentiful Grassfields examples, but the reconstruction PGr *bab`-li is based on class 5 reflexes like Mankon ni-babi-na (see Elias 1984: 39), where the class marker could be a source for final nasals in other groups.

Although the simple inherited velar nasal is almost entirely lost in Grassfields languages, they regularly have the velar nasal as a reflex of PBC * $\eta q \sim PB$ * ηq , e.g. PB * $g \circ \eta q \acute{a}$ 'spear' \sim Bandjoun (Bamileke) kùη, Aghem (Ring) īyōη. Thus, Central Ring forms like Babanki mònlyún 'blood' confirm that PGr inherited the *ng seen in Bantu PB *dóngó 6. The class 6a prefix in Babanki may also explain how a version with a *-ma suffix could generate the many forms in -m that led to the reconstruction of PGr *lém` 'blood' (cf. also Noni (Beboid) ¿lèmέ/bilém 7/8). The presence of the two versions with and without -m in adjacent Momo languages like Njen àluám` and Lower Mundani àlán 7 makes it unlikely that two different roots are involved, but cf. the alternative root PUC *dè 'blood'. Likewise, PGr *tón 'throat' is reflected by velar nasals in every branch of Grassfields and makes it clear that the PBC reconstruction should be PBC *tongo 'throat/voice' (the Isu compound $t \hat{j} \eta \hat{j} \hat{j} \eta$ helpfully seems to preserve some form of V_{2}).

4 Bantu data

Finally, we reach the Bantu languages, which have regularly lost inherited velar nasals, except in relation to a pre-nasalized velar * ηg . Noting the regular correspondence between stem-medial and stem-final * η in Proto-Upper Cross (PUC) and \emptyset in Proto-Bantu, Dim-

mendaal (1978: 233–34) cited $-t\bar{\jmath}\eta$ 'ear' and $-ky\grave{\varrho}\eta$ 'egg' in Nyokon (A45) as exceptional and concluded, "The absence of */ η / in PB can thus easily be explained as an innovation. This innovation could be used for sub-classification (i.e. as a criterion for defining a (very large) subgroup within Broad Bantu). The velar nasal has been lost, e.g., in all but one language of group A 40 in Bantu; / η / is still found in [Nyokon] (A 45) [...]. If the assumption made above is correct, [Nyokon] did not share the innovation and should be reclassified." While Nyokon's phylogenetic placement in the Mbam group (A44–46, A601, A62) is currently not in doubt, the limited number of Bantu languages and words with inherited velar nasals is noteworthy.²⁸

But first, it is important to remember that simple $/\eta/$ is a standard phoneme in the Mbam group today as a regular reflex of PB * ηg , along with occasional $/\eta g/$, as seen in (15).

(15) Examples of Mbam η as the reflex of PB * ηg

'blood' (PB *dúŋgó 6): Nyokon (A45) mànoŋ, Tunen (A44) mànɔŋ, Nomaande (A46) manɔ́ŋɔ́, Tuotomb (A461) mànòŋàm, Nugunu (A622) manɔ́ŋɔ́

'horn' (PB *tóŋgu 5, PGr *ndóŋ): Nyokon (A45) à-ndòom, Tunen (A44) èndóŋ, Nomaande (A46) ɔndóŋó, Tuki (A601) itóngó

4.1 Bantu forms of 'ear'

Now in searching for relics of the inherited simple velar nasal, let us consider the critical word 'ear' (PBC * $tu\eta i$ – PB * $t\acute{o}i$) in Bantu zones AB + Jarawan (6–7). The preservation of the velar nasal is seen in only a few examples (16), with various results otherwise (17).

- (16) Preservation of velar nasal is very rare in Bantu AB, Jarawan
 - a. Nyokon ù-tớŋ and Tuotomb ɔ̂-tớŋà-lɔ̂y (both cl. 3), Yambeta (A462) var. ờ?-tớìŋ
 - b. But no C₂ in Tunen mùlu, Nomande oo-tú, etc.
 - c. Jarawan: Bile $kiiru\eta$, but no C_2 in eight other varieties of Jarawan and A50
 - d. other nasal in Barombi-Kang (A41) dǐwón, Abo (A42) ǐ-wón

²⁸ This Mbam subset has been identified for some time. Most recently, the phylogenies of Grollemund et al. (2015) and Koile et al. (2022: Figures SS1 and SS3) have these same languages in basically similar groupings. The forms cited for the languages in this section come from the database accompanying Grollemund et al. (2015). Tuotomb (Bonek) forms are cited from Mous & Breedveld (1986), which was also a source for some languages in Grollemund.

- (17) When V₂ is preserved, hiatus resolution is frequent
 - a. occasional hiatus, e.g. Lefo (A141) $\hat{\varepsilon}$ -túì
 - b. or glide formation, e.g. Milenge (A15b) *è-tŵ̂*, cf. Koyo (C24) *ì-twéè*
 - c. often y, e.g. Mpongwe (B11a) o-royi, Pinzi (B304) toyi
- d. sometimes y > z, e.g. Punu (B43) di-tudzi, cf. Lega (D25) i-túzi The velar nasal in this root is seen only in Nyokon, Tuotomb, Yambeta and Bile (Jarawan) with some possible remnant nasal features in A41–2 (which are not closely related to Nyokon). Tuotomb clearly has a suffixed form and possibly Nyokon and Yambeta once did too. I do not know enough about Jarawan word formation to speculate on suffixation there, but I note the varieties in reflexes of PR *táònà

I do not know enough about Jarawan word formation to speculate on suffixation there, but I note the variation in reflexes of PB *táànò 'five' in Jarawan: Bile tóŋnó, Kulung túŋnúŋ, Duguri tóón; as well as the coda in Bile murùŋ 'head' (PB *túè).

Nyokon is adjacent to the Eastern Grassfields languages (cf. Bazou tòŋú 'ear'), so one could imagine contact as an explanation, just as

Nyokon is adjacent to the Eastern Grasshelds languages (cf. Bazou tòŋú 'ear'), so one could imagine contact as an explanation, just as Yambeta (A462) nê-bòm 'egg' suggests influence of the distinctive PGr *bùm' 'egg' on a neighboring Bantu language. But the location of Bile in Nigeria would need another explanation.

4.2 Other words in Bantu

Other than in 'ear', the Mbam-Bubi-Jarawan languages lack much evidence for a velar nasal in other words where it would be expected if the languages preserved the inherited phoneme (18).

(18) Bantu forms of other words with inherited velar nasal 'ashes' (PBC *toŋ – PB *túé 3)

A40, 60: Nyokon mù-ól, Nomande mɔotá, Tuki (A601) utú, Tuotomb mɔ̀ɔtɔ́m 6

Bubi obo-tóm

Jarawan: Bile túbú, Duguri túb, Bwazza tú

Although the word for 'ashes' has a similar phonological shape to 'ear', it does not yield a velar nasal in Nyokon or related languages, even when protected by a suffix in several Jarawan languages and Ngoro-Bisoo (A61) li-sip – the suffix is apparently the cl. 14 marker *bu seen as a prefix in Bubi. The final -m in Tuotomb and Bubi looks like the remnant of a class suffix. However, the reduplication in the unusual Lwel (B862) ntip-mtin 'ashes' seems to preserve the original velar nasal. Lwel ntip 'person' shows that initial velar nasal is a typical reflex of the class marker *mtip0 in that language, but in the middle of the word both the original *ntip1 and class prefix ntip2 are preserved (and then the reduplicant copies the coda of the first stem). See also the discussion of Kande (B32) ntip3 and class' above.

'knee' (PCC *dúŋí – PB *dúí 5/6)

Nugunu (A622) i-núu

Bubi (A31) e-ru, Bafia (A53) rì-dú`

Jarawan: Kulung kúlúŋ, cf. Bile nkúŋnú, Bwazza nkúúnù, Duguri kuŋul

Kulung has a velar nasal here, and Duguri perhaps with metathesis. For the Jarawan group, one might reconstruct a protoform with the prefixing and suffixing of the class 15 marker *ku, to which a nasal prefix was added later.

'egg' (PBC *kiŋ – PB *gí 5/6)

Nyokon *nì-kyèn*, Tunen *yòn*, Nomaande *yɔɔ́nɔ́* Tuki (A601) *iqa*, Nukalonge (A62a) *nì-kèé*

Jarawan: Bile *nkì*, Kulung *kiì*

As Dimmendaal noted, there is a final nasal in Nyokon ni-kyèn 'egg', but it is not the velar nasal and not restricted to Nyokon. Since the related A44–46 languages share the nasal, this might be a class marker that became affixed to the monosyllabic root. BLR3 offers a main reconstruction PB *gi 5/6 with several variants: *gé, *gijé, *gijí, and *gidi. This last variant (e.g. Tiene (B81) ma-kuli, Mongo (C61) - $kèl\acute{e}$) probably reflects the incorporation of the class 5 concord marker *di, which would also account for the final nasal in the three A44–A46 languages. Parallel developments can be seen in Koshin (Yemne-Kimbi) $ng\acute{e}n\bar{o}/\eta g\bar{o}$ (5/6), and Afi (Tivoid) $\eta gi\bar{n}/\acute{a}kyi$ where the nasal

suffixes are found only in the singular for 'egg'.²⁹ Likewise in Mambiloid, Ndoro has $\eta_1 \hat{\epsilon} n \bar{a}$ rather than the more typical Mbaw $g^y \hat{\epsilon}/b \hat{\sigma} \eta g^y \hat{\epsilon}$. At this point, we can re-examine the curious final vowel in the PUC reconstruction * $kk\hat{\epsilon}\eta\hat{i}$. PB * $g\hat{i}$ suggests that PUC also had a single syllable root with the class suffix (i.e. * $kk\hat{\epsilon}-n\hat{i}$). The other reconstructed PB variants look like they might be reduplications or contain the class 5 nominal marker *i, hence Vove (B305) $ek\epsilon yi$, but it is possible that a velar nasal developed (see below) and then was lost, e.g. * $kk\hat{\epsilon}-n\hat{i} > kk\hat{\epsilon}-n\hat{i} > kk\hat{\epsilon}-n\hat{i}$ > * $kk\hat{\epsilon}$, cf. PB * $g\hat{i}\hat{i}$.

'mouth, lip' (PBC *nuŋa (ú/ti) – PB *nòà 12, *nyòà)

No velar nasal is observed in the expected languages (Nyokon *nùúl*, Tunen *mùnu*, Yambeta *ù-nù?*, Kulung *kûn*), but frequently there are suffixed class markers in that zone: Tuki *ungúté*, Nugunu *-nyuudé*. Final velar nasals are seen in Mangisa (A63) *ànùŋ*, Eton (A71) *à-nùŋ*, but note the suffix in the related Ewondo (A72a) *anyu-me*.

'louse' (PUC *dán, PLC *lán – PB *dá 9/10)

The most likely case for a velar nasal preservation would be Yambeta ∂n -naŋ, but not in Tunen yìnə or Nomaande weené. Since PB *d > n in these languages, there are many forms with multiple nasals like Ngoro-Asom (A61) npìín or Libie (A62C) ∂n inì, but it becomes difficult to sort them out when one considers the frequency of reduplication for the names of insects, as well as class suffixes.

In general, Bantu looks like Grassfields: there is evidence for the preservation of the velar nasal in the word for 'ear' in a few languages, but otherwise perhaps only in marginal cases like a reduplicated or suffixed form.

5 Analysis

The evidence of the Cross River, Ekoid-Mbe and Bantu languages seems sufficient to reconstruct the distinction between simple $*\eta$

²⁹ I omit Nchanti (Beboid) $g\acute{m}\bar{e}/\bar{a}g\acute{t}$ because Beboid languages show a more general rule for loss of C₂ in class 6 that might be at issue here, e.g. Nchanti $t\bar{e}d\bar{e}/\bar{a}t\bar{a}$ 'stone', Bebe-Jatto $g\acute{a}\beta\acute{t}/g\acute{e}$ 'egg', Akweto $g\acute{t}k\bar{t}/g\acute{t}$ 'egg', $l\bar{s}i/l\acute{t}$ 'eye' (Hombert 1980: 90).

and complex * ηg at the level of Proto-Cross-Congo. But, after several pages of scrutiny, we have found only a few surviving examples of the original simple velar nasal in Bantoid languages. In many groups, we can probably only reconstruct a simple velar nasal in 'ear' and 'ashes', and possibly also 'knee'. In Grassfields, that list is narrowed to 'ear', and in Tivoid and Bantu even that is very marginal. So, among Bantoid branches, Bantu is not really unusual for losing the PCC simple * η , but rather it is unusual for preserving PCC * ηg , which is often reduced to η in most other languages.

A number of questions arise from this review which we will now examine:

- i. In Grassfields and Bantu, we have enough evidence of V_2 to see the regular loss of the velar nasal in C_2 . So, how do we explain the surviving relics?
- ii. Was the loss of the velar nasal independent in each branch or shared by two or more branches?
- iii. Why are there so few words with this phoneme at the PCC stage?

5.1 Explaining the phonological change

Because most Bantoid languages developed secondary velar nasals from original *ŋg or consonants in coda position, velar nasals themselves are quite common in the Bantoid area and their occasional loss is also well known. This loss is quite common in the Teke group of Bantu. For example, in Tiene (B81), NC combinations were simplified to just N, e.g. tùùmà 'cook' (PB *tùmb) and kúóna 'desire' (PB *kúnd-), but the expected velar nasal in túa 'build' (PB *túŋg) was lost (Hyman 2003: 50). The same NC simplifications and loss is seen in Bwala (B70z) búò 'knee' (PB *bóŋgó), ŋkû 'neck' (PB *kíŋgó) (Bollaert et al. 2021). Even in Upper Cross languages, where the velar nasal is generally stable, it was lost in nouns in KoHumono -ttō 'ear', -dū 'knee', -ttò 'road' (PUC *ttòŋ), "but for an unknown reason it was retained in verbs" (Dimmendaal 1978: 97, 105).

Often the deletion leads to assimilation or contraction of the adjacent vowels across morpheme boundaries. In Noni, we saw that the class 5 suffix -e creates an intervocalic environment that deletes stem final g (from *g) with subsequent vocalic assimilation, e.g. * $l\bar{e}g$ - \dot{e} > $l\bar{e}\dot{e}$ (pl. $\bar{e}l\bar{o}\dot{g}$) 'pumpkin' (PB $d\dot{e}g\dot{e}$ 5/6) and * $g\acute{o}g$ - \bar{e} > $g\acute{o}\bar{o}$ (pl. $\bar{e}g\acute{o}g$) 'spear' (PB * $g\grave{o}gg\acute{a}$) (Hombert 1980: 90). In Akoose (A15c), this dele-

tion affected cl. 1 nouns where the suffix then merged with the stem vowel, e.g. Akoose kog 'keep' > nkoo 'keeper' (Hedinger 2008: 6, 26); cf. agents in Babanki (Ring) which show loss and vowel raising, e.g. δ -tag 'stay' > wu-to 'one who stays' (Mutaka-Chie 2006: 84). In both of these languages, the deletion of the velar nasal with vowel effects also affected other contexts including possessive phrases, as seen in Babanki, e.g. ∂sog 'tooth' vs. ∂sog 'dishes' vs. ∂kog 'tooth' vs. ∂sog 'dishes of a person' ∂sog ∂sog ' ∂sog (Akumbu 2016). In Akoose, this same change is still in process, as the velar nasal g frequently elides in a possessive phrase, e.g. ∂sog ∂sog 'tooth (tusk) of elephant'.

In fact, the loss may be less in need of explanation than the preservation is. Why are there relics at all if the loss of the simple velar nasal was a regular phonological change in many Bantoid languages? The best answer seems to be that the velar nasal survived in cases where there was a suffixed class marker. This is perhaps easiest to see in Grassfields, where the PGr reconstructions include a number of stems which incorporated the class 5 concord marker *li as a suffix. Among them is PGr *túŋ-li 'ear' with descendants like Kom àtúŋlá and Mankon atónná. The key feature is that these forms have lost V_o of the root, leaving the velar nasal part of a cluster – presumably that was the phonological environment that blocked its loss. So, the loss of the velar nasal occurred in intervocalic positions, but not in consonant clusters which were generally formed either by suffixes or reduplication. Hence the singular-plural contrast of Evand (Tivoid) k^{w} ótóngók/átō based on the difference between the consonantal cl. 15 *ku and vocalic cl. 6 *a suffixes.

Because of the uncertainty in reconstructing many velar nasals in final position, it is not clear whether final position was regularly an environment for the loss. Instances where the inherited velar nasal is now found intervocalically and finally are presumably later developments, i.e. Bazou $t \delta \eta u$ arose from a later simplification of the * ηl cluster and Bandjoun $t \delta \eta$ from the eventual loss of the final vowel of the suffix as well. Since the two most common conditions needed for the original velar nasal's preservation are class marker suffixation and loss of V_2 , and both of those conditions are rare in Bantu outside zone A, it is not surprising that the rest of Bantu has almost no examples of velar nasal preservation.

The suffixation of concord markers was a variable process, due to individual lexemes and the frequency of various syntactic structures. For example, we might reconstruct forms of 'ear' in two classes, which probably occurred both in Grassfields and Bantu languages:

Pr-Bantu-Grassfields *ku- $tu\eta i$ (15/6) > * $kut\acute{v}i$ [η loss] (the common pattern in Bantu) Pr-Bantu-Grassfields * $tu\eta i$ -li (5/6) > * $tu\eta li$ [vowel loss] (the common pattern in Grassfields)

In languages like the Grassfields group, which lost class 15, it is not surprising that class 5 was used for the singular of class 6. Presumably the class 5 concord marker *li became re-analyzed as a suffix and frozen in certain contexts leading to allomorphy with the plural or remnant *ku- forms. The difference between these variants was originally morphological and then would have become phonological through the relevant losses, with the allomorphs being subject to analogical processes of levelling.

5.2 Was this loss independent in the various Bantoid groups?

The losses of the PCC simple velar nasal must have happened before the merger of * η and * ηg (or otherwise both would be lost), so we might posit the following processes of development (which did not happen in all languages):

Initial conditions of PCC: distinct * η and * ηg

- 1. cluster formation (via suffixation, reduplication, V_2 loss) ongoing process
- 2. loss of * η except in cluster environments
- 3. merger of * η and * $\eta g > *\eta$

Ekoid alone seems to preserve inherited velar nasals so it must have engaged only in Process 3 without any loss of * η , and Mbe engaged in Process 2 without apparent cluster formation. So, we can put aside Ekoid-Mbe from any developments shared with others. On the other hand, Grassfields and Bantu languages are closely related and have similar results (no inherited velar nasals except in 'ear'), so it is likely that their loss happened at some stage common to them. But most of the other Bantoid branches had significant exposure to the suffixation of class markers, V_2 loss and * $\eta g > *\eta$ in ways that could be independent or areal, so the loss of * η may also have been independent. The limited number of relevant roots reconstructed at the

PCC stage and the limited number of exceptions makes it hard to know what a general pattern is.

5.2 The origin of * η

The reconstructed Proto-Benue-Congo velar nasal has an uneven distribution: it is never initial, and even in $\rm C_2$ position it is not very common. It is true that PBC reconstructions are at an early stage and mostly just exist for nouns, but the marginality of the velar nasal suggests that it might have developed through conditioning in a narrow environment. For a clue to the possible environment, let us look at the word for 'firewood'. In Bantu we see doublets, one of which curiously shows the loss of a medial nasal:

PB *kúnì (9/10, 11/10) zones ABCD+ (BLR 2042)

PB *kύὶ (9/10, 11/10) zones BCEF+ (BLR 1983)

Usually, a C_2 nasal is one of the most stable sounds in Bantu languages, but we could explain this loss if the nasal had become a velar nasal in some environment: $*n > *y > \emptyset$. In the case of 'firewood', the nasal's environment is between high back and front vowels, which is exactly the same environment we see in many of the words reconstructed with velar nasals: PBC *tuŋi 'ear', PCC *tóŋé 'ashes', PUC *dúŋí 'knee', PUC dáŋ (Sur dani) 'louse'.³¹ Did those velar nasals also develop from earlier alveolar or dental nasals? Was Mukarovsky right in reconstructing early *n in some words?

If we look at de Wolf's PBC reconstructions for 'wood', 'fire' and 'firewood', we see several related forms which look like they have a common stem but vary by class and the addition of a suffix -i.³²

PBC *kwon (*ka/*ti) 'tree'

PBC *kwoni (*bu/*î) 'firewood' – cf. BCCW #kónì

PBC *zwuŋi (*ku/*a) 'fire' – cf. PUC *k"ón 'fire'

We might want to reconstruct just one root *kón with 'tree/wood' as the basic meaning but specified semantically by different class markers. In several Tiv and Grassfields languages, one class is used for 'tree' and another for 'firewood'. The suffix -i looks like the con-

³⁰ The restriction of velar nasals to non-initial position is fairly common, occurring in about one-third of the languages which have the phoneme (Anderson 2013).

³¹ This environment was operative in Umbundu (R11), where we see the loss of a nasal after a back vowel and before i not only in $\acute{o}l\acute{o}-hw\acute{t}$ 'firewood' but also in $\acute{e}-kw\acute{t}$ 'ten' (PB $*k\acute{o}m\grave{t}$) and $\acute{o}-s\acute{o}y\grave{t}$ 'shame' (PB $*c\acute{o}n\grave{t}$).

³² For fuller data on this root, see Williamson (1989: 250-53).

cord marker for classes 9 and 10 – exactly the gender of PB *kónì 'firewood'. So, these various allomorphs would provide a way to explain the presence or absence of the necessary conditioning environment for producing a velar nasal.

Another explanation for the restricted development of the velar nasal might be to start from * ηg , just as final / η / in modern English arose from / ηg / after the loss of final /g/, as the spelling in spelling still indicates. Perhaps the incorporation of class suffixes (perhaps with syncope?) created a cluster which led to the deletion of *g: e.g. * $tu\eta g(i)$ -di 'ear' > * $tu\eta g$ -li > PGr * $tu\eta$ -li, just as / ηg / is sometime reduced to / η / in pronunciations of the medial clusters in language and English. We might also wonder whether * ηg could also sometimes be reduced by a following *i, in the pattern above?

Proto-Bantu has some, although not many, verb roots with double vowels or diphthongs which could in theory be the results of a consonant loss. As far as I know, none has been proposed as the result of the loss of the velar nasal – is that because PBC verbs were CVC and none incorporated a high-vowel suffix of the sort needed to generate the velar nasal?

6 Conclusion

It is not clear how far back the simple velar nasal $*\eta$ itself should be reconstructed, but there is good reason to reconstruct five or more Proto-Cross-Congo noun roots containing it. However, its marginal status made it easy to be lost or merged with $*\eta g$. The Ekoid-Mbe branch is distinctive in that it preserved the simple velar nasal in at least five of the stems seen in PCC, so its subsequent loss in Mbe is a phonological innovation. There are fewer surviving examples in other Bantoid languages, mostly just in the words 'ear' and 'ashes', often in suffixed forms. There is not enough data to support grouping together Bantoid branches based on this criterion, except possibly Grassfields with Bantu. The Grassfields examples particularly provide evidence to think that exemption from loss could be triggered by an environment in a consonant cluster caused by V_2 loss before a suffix.

Particularly useful is the fact that some Bantoid languages (most clearly Mbe) seem to parallel Bantu in having a distinction in reflexes of reconstructed PBC * ηq and * η . This parallel confirms that the dis-

tinction should be reconstructed at the PCC stage and also enables improvements for the PBC stage. For example, contrary to de Wolf, I would reconstruct PBC *ŋg in the words for 'blood' (de Wolf *luŋ), 'horn, tusk' (de Wolf *tano), and possibly 'leopard' (de Wolf *kpoŋi). The absence of final velar nasals in the Bantoid forms for certain words also suggests that velar nasals should be removed from the reconstructions of the roots for 'mosquito, bee', 'egg', and 'wing'.

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Abbreviations

BCCW = Benue-Congo Comparative Wordlist

BCR = Benue-Congo Reconstructions

BLR = Bantu Lexical Reconstructions

C = consonant

GBWG = Grassfields Bantu Working Group

NC = Niger-Congo

PB = Proto-Bantu

PBC = Proto-Benue-Congo

PCC = Proto-Cross-Congo

PEM = Proto-Ekoid-Mbe

PGr = Proto-Grassfields

PJ = Proto-Jukunoid

 $PLC = Proto-Lower\ Cross$

PP = Proto-Plateau

 $PUC \,=\, Proto\text{-}Upper\ Cross$

PWN = Proto-Western Nigritic (Mukarovsky)

V = vowel

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