Calculation of the C.I.P.W. norm

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Summary. A modification of the rules for the calculation of C.I.P.W. norms when no normative quartz is present is suggested.

IN the course of writing a programme to calculate C.I.P.W. norms on EDSAC II, it became apparent that the rules, as given by Washington (1917), Johannsen (1939), and other authors, for calculating the norm in the absence of normative quartz were inconvenient. Modification of the method hitherto in common usage produces a calculation of greater simplicity.

Normative quartz is present in a great many igneous rocks and it is therefore usual to assume that quartz will occur in the norm and to form, where appropriate, the provisional normative minerals of highest silica content, orthoclase, albite, diopside, hypersthene or wollastonite, and sphene. Any excess of silica after the allocation of silica necessary to form the minerals of the norm is calculated as quartz. If, however, there is insufficient silica in the rock to combine with all other relevant oxides to give the highly silicic minerals, then these provisional minerals must be converted to alternative minerals of lower silica content.

The provisional normative minerals are listed in table I in the order in which they must be converted to alternative minerals of lower silica content. Each mineral in turn is changed to the alternative subsilicic mineral until the silica accommodated in the normative minerals is equal to the silica content of the rock. This redistribution of silica can be done quickly by considering the silica deficiency defined as the amount¹ of silica required for the formation of the normative minerals so far formed, less the amount of silica in the rock, at each stage of the redistribution. Suppose that *one* molecule of a given provisional mineral yields Umolecules of silica and V molecules of the corresponding mineral of lower silica content. In this context U and V need not necessarily be

¹ The amount of a given mineral, oxide or element throughout this paper is expressed in terms of its molecular proportion, i.e. the weight percentage divided by the formula weight.

whole numbers; they can be simple fractions. If exactly one molecule of silica is formed by conversion of the provisional mineral then the number of molecules of the alternative mineral produced will be W = V/U. Thus, if we have an amount X of the provisional mineral, then it can yield a maximum amount XU of silica. Suppose that at a given

TABLE I

A_1	A_2	B_1	B_2	U	V	V	V
hypersthene sphene albite orthoclase wollastonite	olivine perovskite nepheline leucite dicalcium silicato	$\begin{array}{l} ({\rm Mg},{\rm Fe}){\rm O}.{\rm SiO}_{2} \\ {\rm CaO}.{\rm TiO}_{2}.{\rm SiO}_{2} \\ {\rm Na}_{2}{\rm O}.{\rm Al}_{2}{\rm O}_{3}.6{\rm SiO}_{2} \\ {\rm K}_{2}{\rm O}.{\rm Al}_{2}{\rm O}_{3}.6{\rm SiO}_{2} \\ {\rm CaO}.{\rm SiO}_{2} \end{array}$	$2(Mg, Fe)O.SiO_2$ CaO.TiO_2 Na ₂ O.Al ₂ O ₃ .2SiO ₂ K ₂ O.Al ₂ O ₃ .4SiO ₂ 2CaO.SiO ₂	$\frac{1}{2}$ 1 4 2 $\frac{1}{2}$			1114121
diopside leucite	dicalcium silicate olivine kaliophilite	$ \left. \begin{array}{l} {\rm CaO.(Mg,Fe)O.2SiO}_2 \\ {\rm K_2O.Al_2O_3.4SiO}_2 \end{array} \right. $	$\left\{\begin{array}{l} 2\mathrm{CaO.SiO}_2\\ \\ 2(\mathrm{Mg,Fe)O.SiO}_2\\ \mathrm{K}_2\mathrm{O.Al}_2\mathrm{O}_3.2\mathrm{SiO}_2 \end{array}\right\}$	1 2	$ \left\{\begin{array}{c} \frac{1}{2} \\ \frac{1}{2} \\ \frac{1}{2} \\ 1 \end{array}\right. $		12 1212

 A_1 . Name of provisional mineral.

A₂. Name of alternative subsilicic mineral.

 B_1 . Formula of provisional mineral.

 B_2 . Formula of alternative subsilicic mineral.

U. Number of molecules of silica formed when one molecule of the provisional mineral is converted.

V. Number of molecules of alternative subsilicic mineral formed from one molecule of provisional mineral.

W. Number of molecules of alternative mineral formed when one molecule of silica is formed from the provisional mineral.

stage of the calculation, this provisional mineral is the next to be converted to its alternative and that the silica deficiency at this stage of the calculation is D. Then if D > XU, all the provisional mineral must be converted to the alternative, forming an amount equal to XVof the subsilicic mineral and releasing an amount XU of silica. This reduces the deficiency to (D-XU). If however D < XU then it is unnecessary to change the whole amount of the provisional mineral into the alternative subsilicic mineral; the deficiency is eliminated by conversion of an amount equal to D/U. Thus an amount (X-D/U) of the provisional mineral remains in the norm and an amount WD of the subsilicic mineral is formed. The values of U, V, and W for each provisional mineral are set down in table I. In redistributing silica in the absence of normative quartz the silica deficiency is calculated and the provisional minerals converted to the corresponding subsilicic minerals in the manner indicated until the deficiency has been annulled.

This method of calculating the C.I.P.W. norm is incorporated in the following rules for calculating the norm. The list of possible normative minerals with their formulae adopted for the purposes of calculation of the norm are set down in table II. In this set of rules the abbreviation for a normative mineral is taken to mean the amount of that mineral present in the norm expressed as its molecular proportion. If the abbreviation is primed then it denotes the amount of a normative

TABLE II. Normative mineral molecules

Quartz (Q)	SiO,
Corundum (C)	Al ₂ Õ ₂
Zircon (Z)	ZrO, SiO,
Orthoclase (or)	K.O.Al.O.68iO.
Albite (ab)	Na.O.Al.O.68iO.
Anorthite (an)	CaO.Al,O.2SiO,
Leucite (lc)	K ₂ O.Al ₂ O ₃ .4SiO ₂
Nepheline (ne)	Na ₂ O.Al ₂ O ₃ .2SiO ₂
Kaliophilite (kp)	K20.Al202.2SiO2
Halite (hl)	NaCl
Thenardite (th)	Na ₂ O.SO ₃
Sodium carbonate (nc)	$Na_2O.CO_2$
	N OF O INO
Acmite (ac)	$Na_2O.Fe_2O_3.4SiO_2$
Sodium metasilicate (ns)	$Na_2 O.SiO_2$
Potassium metasilicate (ks)	$K_2 O.SiO_2$
Diopside (di)	$CaO.(Mg, Fe)O.2SiO_2$
Wollastonite (wo)	CaO.SiO ₂
Hypersthene (hy)	$(Mg, Fe)O.SiO_2$
Olivine (ol)	$2(Mg,Fe)O.SiO_2$
Dicalcium silicate (cs)	2CaO.SiO ₂
Magnetite (mt)	$\mathrm{FeO.Fe}_{2}\mathrm{O}_{3}$
Chromite (cm)	$FeO.Cr_2O_3$
Ilmenite (il)	$FeO.TiO_2$
Hematite (hm)	$\mathrm{Fe}_{2}\mathrm{O}_{3}$
Sphene (tn)	$CaO.TiO_2.SiO_2$
Perovskite (pf)	CaO.TiO ₂
Rutile (ru)	TiO ₂
Apatite (ap)	$3CaO.P_2O_5.\frac{1}{3}CaF$
Fluorite (fr)	CaF _a
Pyrite (pr)	FeS2
Calcite (cc)	CaO.CO.

mineral which has been formed provisionally. This amount may subsequently be reduced if there is no normative quartz present. The use of an oxide formula or the symbol of an element is taken to indicate the amount of that oxide or element available for the formation of normative minerals at that stage of the calculation. That is to say the total amount of oxide or element present in the rock less the amount of that oxide or element already used in the formation of normative minerals. As each normative mineral is formed the amount of silica, if any, required to form this mineral is added to the amount of silica Y necessary for the formation of the minerals in the norm.

The revised rules proposed for the calculation of the norm are:

- 1. Calculate the amounts (molecular proportions) of the oxides and elements present by dividing the weight percentages by the appropriate formula weights.
- 2. Add (MnO + NiO) to FeO.
- 3. Add (BaO + SrO) to CaO.
- 4. Set $Z = \operatorname{ZrO}_2$; set Y = Z.
- 5. Set $ap = P_2O_5$; subtract an amount equal to $\frac{10}{3}ap$ from CaO. If fluorine is present and if $F > \frac{2}{3}ap$ subtract an amount equal to $\frac{2}{3}ap$ from F. If $F < \frac{2}{3}ap$ all the fluorine is allocated to ap.
- 6. Set fr = F/2; subtract an amount equal to fr from CaO.
- 7. Set hl = Cl; subtract an amount equal to hl/2 from Na₂O.
- 8. If SO₃ is to be calculated as SO₃ and not as S, then set $th = SO_3$; subtract an amount equal to th from Na₂O.
- 9. If S is present or if SO₃ is to be calculated as S then put pr = S/2 or SO₃/2; subtract an amount equal to pr from FeO.
- 10. (a) If the rock contains cancrinite then put $nc = CO_2$; subtract an amount equal to nc from Na₂O.
 - (b) If the rock contains calcite set $cc = CO_2$; subtract an amount equal to cc from CaO. If the modal calcite is secondary or from associated limestone it is not included in the norm.
- 11. Set $cm = Cr_2O_3$; subtract an amount equal to cm from FeO.
- 12. If $\text{FeO} > \text{TiO}_2$ set $il = \text{TiO}_2$ and subtract an amount equal to il from FeO. There is no further TiO_2 available to form other normative minerals.
 - If FeO < TiO₂ set il = FeO and subtract an amount equal to il from TiO₂. There is no further FeO available to form other normative minerals.
- 13. If $Al_2O_3 > K_2O$ set $or' = K_2O$ and subtract an amount equal to or' from Al_2O_3 ; add an amount equal to 6or' to Y.
 - If $Al_2O_3 < K_2O$ set $or' = Al_2O_3$ and subtract an amount equal to or' from K_2O . There is no further Al_2O_3 available to form other normative minerals. Set $K_2O = ks$; add an amount equal to (6or' + ks) to Y.
- 14. If $Al_2O_3 > Na_2O$ set $ab' = Na_2O$ and subtract an amount equal to ab' from Al_2O_3 . There is no further Na_2O available to form other normative minerals. Add an amount equal to 6ab' to Y.

C. H. KELSEY ON

- If $Al_2O_3 < Na_2O$ set $ab' = Al_2O_3$ and subtract an amount equal to ab' from Na₂O. There is no further Al_2O_3 available to form other normative minerals. Add an amount equal to 6ab' to Y.
- 15. If $Na_2O > Fe_2O_3$ set $ac = Fe_2O_3$ and subtract an amount equal to ac from Na_2O . There is no further Fe_2O_3 available to form other normative minerals. Set $ns = Na_2O$; add an amount equal to (4ac+ns) to Y.
 - If $Na_2O < Fe_2O_3$ set $ac = Na_2O$; subtract an amount equal to ac from Fe_2O_3 ; add an amount equal to 4ac to Y.
- 16. If $Al_2O_3 > CaO$ set an = CaO and subtract an amount equal to an from Al_2O_3 . There is no further CaO available to form other normative minerals. Add an amount equal to 2an to Y; set $C = Al_2O_3$.
 - If $Al_2O_3 < CaO$ set $an = Al_2O_3$; subtract an amount equal to an from CaO; add an amount equal to 2an to Y.
- 17. If $\text{CaO} > \text{TiO}_2$ set $tn' = \text{TiO}_2$; subtract an amount equal to tn' from CaO; add an amount equal to tn' to Y.
 - If CaO < TiO₂ set tn' = CaO and subtract an amount equal to tn' from TiO₂. There is no further CaO available to form other normative minerals. Set ru = TiO₂; add an amount equal to tn' to Y.
- 18. If $\operatorname{Fe_2O_3} > \operatorname{FeO}$ set $mt = \operatorname{FeO}$ and subtract an amount equal to mt from $\operatorname{Fe_2O_3}$. There is no further FeO available to form other normative minerals. Set $hm = \operatorname{Fe_2O_3}$.
 - If $\text{Fe}_2O_3 < \text{FeO}$ set $mt = \text{Fe}_2O_3$; subtract an amount equal to mt from FeO.
- 19. Add MgO to FeO to form (Mg,Fe)O. Calculate the ratios MgO/ (MgO+FeO) and FeO/(MgO+FeO) and use these relative proportions to calculate the weight percentages of diopside, hypersthene, and olivine.
- 20. If CaO > (Mg,Fe)O set di' = (Mg,Fe)O: subtract an amount equal to di' from CaO; set wo' = CaO; add an amount equal to (2di' + wo') to Y.
 - If CaO < (Mg,Fe)O set di' = CaO; subtract an amount equal to di' from (Mg,Fe)O; set hy' = (Mg,Fe)O; add an amount equal to (2di'+hy') to Y.
- 21. Y now gives the amount of silica required for all the normative minerals so far formed.

If $SiO_2 > Y$ set $Q = SiO_2 - Y$. The calculation of the norm is

280

completed by conversion of the molecular proportions to weight percentages of normative minerals.

If $\operatorname{SiO}_2 < Y$ the deficiency $D = Y - \operatorname{SiO}_2$ and further calculations have to be made (rules 22-27). These calculations must be continued until the deficiency has been reduced to zero. The calculation is then completed by conversion of the molecular proportions to weight percentages of the normative minerals.

TABLE III. Calculation of the C.I.P.W norm of a bostonite from Maros, Celebes

				No	rmative					
Wt. %	Amount			m	inerals	Y				
61.15	1.019			il	0.003					
22.07	0.217, 0.143, 0	048, —		or'	0.074	0.444				
1.05	0.006,			ab'	0.095	0.570				
1.02	0.014, 0.011, 0)·005,] _{0.1}	015	an	0.014	0.028				
0.40	0.010,	}0.	015,	C	0.034					
0.75	0.014,			mt	0.006					
5.86	0·095, —			hy'	0.012	0.012				
7.01	0.074,									
0.71						1.057				
0.21	0.003,									
$\frac{\text{MgO}}{(\text{Mg,Fe})\text{O}} = \frac{2}{3}; \qquad \frac{\text{FeO}}{(\text{Mg,Fe})\text{O}} = \frac{1}{3}.$ $D = 0.038, \ hu'/2 = 0.008 \therefore \ ol = 0.008; \ hu = 0; \ D_{2} = 0.030.$										
4ab	$' = 0.380$ \therefore at	b = 0.087	; $ne = 0$	·00ŝ.						
Norm										
С	3.47	ol	1.21							
01	41.14	\mathbf{mt}	1.39							
al	o 45.59	il	0.46							
aı	1 3 ·89									
ne	e 2·27									
	Wt. % 61.15 22.07 1.05 1.02 0.40 0.75 5.86 7.01 0.71 0.21 $\overline{(1)}$ 4ab C C on all arr no	Wt. % Amount 61.15 1.019 22.07 0.217, 0.143, 0 1.05 0.006, 1.02 0.014, 0.011, 0 0.40 0.010, 0.75 0.014, 5.86 0.095, 7.01 0.074, 0.21 0.003, $\frac{MgO}{(Mg,Fe)O} = \frac{2}{3};$ 4.0.038. $hy'/2 = 0.008 \therefore old 4ab' = 0.380 \therefore all No C 3.477 or 41.14 ab 45.59 an 3.889 ne 2.27$	Wt. % Amount 61.15 1.019 22.07 0.217, 0.143, 0.048, — 1.05 0.006, — 1.02 0.014, 0.011, 0.005, 0.040 0.40 0.010, 0.075 , 0.014 , — 5.86 0.095, — 7.01 0.074, — 0.21 0.003, — $\frac{MgO}{(Mg,Fe)O} = \frac{2}{3}; \frac{FeO}{(Mg,Fe)}$ 4ab' = 0.380 \therefore ab = 0.087 Norm C 3.47 ol or 41.14 mt ab 45.59 il an 3.89 ne 2.27	Wt. % Amount 61.15 1.019 22.07 0.217, 0.143, 0.048, — 1.05 0.006, — 1.02 0.014, 0.011, 0.005, 0.015 , — 0.40 0.010, 0.075 0.014 , — 5.86 0.095, — 7.01 0.074, — 0.21 0.003, — $\frac{MgO}{(Mg,Fe)O} = \frac{2}{3};$ FeO $\frac{FeO}{(Mg,Fe)O} = \frac{1}{3}$. 4ab' = 0.380 \therefore ab = 0.087; ne = 0 Norm C 3.47 ol 1.21 or 41.14 mt 1.39 ab 45.59 il 0.46 an 3.89 ne 2.27	Wt. % Amount m 61·15 1·019 <i>il</i> 22·07 0·217, 0·143, 0·048, — or' 1·05 0·006, — ab' 1·02 0·014, 0·011, 0·005, 0.015 , — an 0·40 0·010, 0.015 , — mt 5·86 0·095, — hy' 7·01 0·074, — 0.71 — 0.71 — 0.21 0·003, — $\frac{MgO}{(Mg,Fe)O} = \frac{2}{3}; \frac{FeO}{(Mg,Fe)O} = \frac{1}{3}.$ 4ab' = 0·380 \therefore ab = 0·087; ne = 0·008. Norm C 3·47 ol 1·21 or 41·14 mt 1·39 ab 45·59 il 0·46 an 3·89 ne 2·27	Wt. % Amount minerals 61·15 1·019 il 0·003 22·07 0·217, 0·143, 0·048, — or' 0·074 1·05 0·006, — ab' 0·095 1·02 0·014, 0·011, 0·005, $0·015$, — an 0·014 0·40 0·010, $0·015$, — ar 0·014 0·75 0·014, — mt 0·006 5·86 0·095, — hy' 0·015 7·01 0·074, — 0·21 0·003, — $\frac{MgO}{(Mg,Fe)O} = \frac{2}{3}; \frac{FeO}{(Mg,Fe)O} = \frac{1}{3}.$ 4 0·038. $hy'/2 = 0.008 \therefore ol = 0.008; hy = 0; D_1 = 0.030.$ $4ab' = 0.380 \therefore ol = 0.08; hy = 0; D_1 = 0.030.$ $Aab' = 0.380 \therefore ol = 0.087; ne = 0.008.$ Norm C 3·47 ol 1·21 or 41·14 mt 1·39 ab 45·59 il 0·46 an 3·89 ne 2·27				

22. If D < hy'/2 set ol = D and hy = hy'-2D. The silica deficiency is now zero.

If D > hy'/2 set ol = hy'/2 and hy = 0; put $D_1 = D - hy'/2$.

23. If $D_1 < tn'$ set $tn = tn' - D_1$ and $pf = D_1$. The silica deficiency is now zero.

If $D_1 > tn'$ set pf = tn' and tn = 0; put $D_2 = D_1 - tn'$.

24. If $D_2 < 4ab'$ set $ne = D_2/4$ and $ab = ab' - D_2/4$. The silica deficiency is now zero.

If $D_2 > 4ab'$ set ne = ab' and ab = 0, put $D_3 = D_2 - 4ab'$.

25. If $D_3 < 2or'$ set $lc = D_3/2$ and $or = or' - D_3/2$. The silica deficiency is now zero.

If $D_3 > 2or'$ set lc' = or' and or = 0; put $D_4 = D_3 - 2or'$.

- 26. If $D_4 < wo'/2$ set $cs = D_4$ and $wo = wo'-2D_4$. The silica deficiency is now zero.
 - If $D_4 > wo'/2$ set cs = wo'/2 and wo = 0; put $D_5 = D_4 wo'/2$.
- 27. If $D_5 < di'$ add an amount equal to $D_5/2$ to the amounts of cs and ol already in the norm; set $di = di' D_5$. The silica deficiency is now zero.
 - If $D_5 > di'$ add an amount equal to di'/2 to the amounts of cs and ol already in the norm; put di = 0 and $D_6 = D_5 di'$.
- 28. Set $kp = D_6/2$ and $lc = lc' D_6/2$. The silica deficiency is now zero.

To illustrate the use of these rules, the C.I.P.W. norm of a bostonite from Maros, Celebes (Washington, 1917, p. 274, analysis no. 37), is calculated in table III.

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References

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