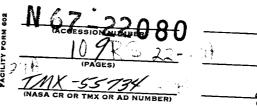
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3 NEWLY IDENTIFIED LINES IN THE Ne I ISOELECTRONIC SEQUENCES 6

U. FELDMAN IL. COHEN



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GODDARD SPACE FLIGHT CENTER
GREENBELT, MARYLAND

NEWLY IDENTIFIED LINES IN THE Ne I ISOELECTRONIC SEQUENCES

Ву

U. Feldman* and L. Cohen Goddard Space Flight Center Greenbelt, Maryland

ABSTRACT

Using a grazing-incidence spectrometer and a low inductance, $14\mu F$, 12-17 kV, spark source, the authors have observed spectra of Sc x11, Ti x111, and V x1V. The lines have been identified as arising from transitions between the ground level $2s^22p^{6-1}S_0$ and the following electronic configurations: $2s^22p^6-3p$, and $2s^2-2p^5-4s$, 4d, 5d. The transitions of the type $2s^2-2p^6-2s^2-2p^5$ 4d have been observed also in Co xV111 Ni x1x and Cu xx.

^{*}NASA-National Academy of Science - National Research Council Postdoctorial Research Associate.

It is generally evident on considering an isoelectronic sequence of ions that as one proceeds to higher degrees of ionization the number of known levels decreases. One of the important reasons for this situation, especially for the very high degrees of ionization (above the 10th, say), is the lack of a suitable source for generating the spectra with reasonable intensity. In the Ne I isoelectronic sequence energy levels have been identified as far as Zn xx1. Edlen and Tyren (1936) treated the following ions: K 1x, Ca x1, Sc xii, Ti xiii, and V xiv. The spectra of Cr xv, Mn xvi, Fe xvii, and Co xviii were reported by Tyren (1938). Fawcett (1965) measured some levels of Sc X11, Ti X111 and V X1V, Feldman, Cohen, and Swartz (1967) extended the isoelectronic sequence to Ni xix, Cu xx, and Zn xxi. Until recently, a noticeable gap in the Ne I isoelectronic sequence existed for the spectra of Sc x11, Ti x111, and V x1v where only the following levels had been identified: $2s^2 2p^{6-1}S_0$, $2s^2 2p^5 3s ^3P_1^{\alpha}$, $^1P_1^{\alpha}$, and $2s^2 2p^5 3d ^3P_1^{\alpha}$, $^3D_1^{\alpha}$, $^1P_1^{\alpha}$ (the $2s^2 2p^5 3d ^3P_1^0$ level in V x1v not being known). For ions below Sc X11, and above V X1V, the number of known levels is usually greater. This gap was probably due to the fact that the experimental methods by which the Sc, Ti and V ions were first investigated (Edlen and Tyren, 1936) were inadequate to obtain high intensities in this region. Therefore, only

the most intense lines were observed. Later when Tyrén (1938) treated the ions from Cr xv to Co xv111 with improved methods and identified a greater number of energy levels, he did not return to the Sc, Ti and V ions.

Fawcett (1965), using a two-meter grazing incidence spectrometer and a 600 lines per mm Bausch and Lomb blazed replica grating, with a $1/2\mu F$, 90 kV vacuum spark source, recorded lines of Sc, Ti and V. He classified three lines each of Sc x11 and Ti x111, and four lines of V x1V belonging to transitions from the following configurations: $2p^6-2s$ $2p^6$ 3p and 2p 6-2p 4d. He also reports an additional line in V x1V classified as follows: 2p 6-2p 3d. His wavelength measurements are reported to an accuracy of $\pm 0.02A$.

Independently, we recorded spectra in this region by using a modified Jarrell-Ash 3-meter grazing-incidence spectrometer, with a Bausch and Lomb 1200 lines per mm blazed gold replica. Our source was a 14µF, 12 to 17 kV, low-inductance condensed spark. As reference for wavelength calibration we used carbon and oxygen lines, and known lines of the same element isoelectronic with Ne I (Moore, 1949, 1952). We have observed spectra of Sc x11, V x111, Ti x1v, Co xv111, Ni x1x and Cu xx. As well as the lines arising from transitions between the 2s 2 2p 6 and 2s 2 2p 5 3s, 3d electronic configurations, the lines arising from transitions between the following configurations

were recorded for Sc X11, V XV111 and Ti X1V: $2s^2 \ 2p^6 - 2s \ 2p^6 \ 3p$, and $2s^2 \ 2p^6 - 2s^2 \ 2p^5 \ 4s$, 4d, 5d. The

wavelengths, energies, and visually estimated intensities

are summarized in Table 1. Term values are given in Table 2.

The line belonging to $2s^2 \ 2p^6 \ So - 2s^2 \ 2p^5 \ 3d \ 3p_1^o$ in V X1V

is also included. The wavelengths in Table 1 were measured

to an accuracy of better then $\pm 0.005A$. For Co XV111 Ni X1X

and Cu XX transitions of the type $2s^2 \ 2p^6 - 2s^2 \ 2p^5 \ 4d$ have

been measured to an accuracy of $\pm 0.01A$. The wavelengths and energies are in Table 3.

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TABLE 1 - Classified lines from the 2s 2 $2p\,^6$ - 2s $2p^6$ 3p and $2s^2 2p^6 - 2s^2 2p^5 3d$, 4s, 4d, 5d transitions of Sc $\boldsymbol{x}\textsc{il}$, Ti $\boldsymbol{x}\textsc{il}$ and V $\boldsymbol{x}\textsc{iv}$

5	Int \circ (cm ⁻¹)	4691000	5299400	5324200	5632500	5689900	5850000	5903500	6350000	6406600
V X1V	Int	10	n	7		~	4	5	7	2
	λ (A)	21.294	18.870	18,783	17.754	17.575	17.094	16,939	15.748	15.609
Н	$c(cm^{-1})$ $\lambda(\mathtt{A})$		4733300	4754000	4966500	5014300	5163700	5207200	5596300	5641100
Ti X111	Int		7	7	Ĺ.	Н	4	5		7
	(A)		21.127	21.035	20.135	19.943	19.366	19.204	17.869	17.727
	(cm^{-1}) (A)		4198000 21.127	4215000	4339300	4378800 19.943	4521000	4557900	4892800	4926600 17.727
Sc X11	Int		H		~	,-i	4	7.7		7
	λ(A)		23.821	23.725	23.045	22,837	22.119	21.940	20,438	20.298
		3 P 0	$^3_{\rm P}$	1 $^{\mathbf{P}_{0}}$	3. Po	$^{1}P_{1}^{0}$	3D_1	1 P ₁	$^3D_1^0$	$^{1}\mathbf{P}_{1}^{0}$
	Transitions	$^{1}\text{S}_{0} - 2\text{s}^{2} 2\text{p}^{5}3\text{d}$	2s 2p ⁶ 3p	$2s^2 2p^5 3p$	$2s^2 2p^5 4s$	$2s^2 2p^5 4s$	$2s^2 2p^5 4d$	$2s^2 2p^5 4d$	$2s^2 2p^55d$	$2s^2 2p^5 5d$
	Tran	$2s^22p^6$ 1S_0	=	Ξ	=	£	=	Ξ	=	Ξ

TABLE 2 - Energy scheme for terms in Sc X11, Ti X111, and V X1V

Configuration	Designation	,	Sc x 11 (cm ⁻¹)	Ti x111 (cm ⁻¹)	$V \times V (cm^{-1})$
2s ² 2p ⁵ 3d	3d 3p0	-			4596100
2s 2p ⁶ 3p	3p 3p ⁰	Н	4198000	4733300	5299400
2s 2p ⁶ 3p	3p ¹ p ⁰	Н	4215000	4754000	5324200
$2s^2 2p^5 4s$	4s 3p0	Н	4339300	4966500	5632500
$2s^2 2p^5 4s$	4 ¹ P ⁰	7	4378800	5014300	5689900
$2s^2 2p^5 4d$	² 0 Pγ	Н	4521000	5163700	5850000
$2s^2 2p^5 4d$	4d ¹ P ⁰	, -	4557900	5207200	5903500
$2s^2 2p^5 5d$	5d 3D0	1	4892800	5596300	6350000
$2s^2 2p^5 5d$	5d ¹ P ⁰	7	4926600	5641100	9406600

TABLE 3 - Classified lines from the $2s^2\ 2p^6 - 2s^2\ 2p^5$ 4d transitions of Co xv111, Ni x1x and Cu xx

	Co	Co xviii	Ni X1X	X1X	Cr	Cu xx
Transitions	λ(A)	$\cup (\operatorname{cm}^{-1}) \lambda(\mathbf{A})$	λ(A)	∪(cm ⁻¹)	λ(Α)	$\lambda(\mathbf{A})$ $\delta(\mathbf{cm}^{-1})$
$2s^2 2p^{6} ^{1}s_0 - 2s^2 2p^5 4d ^{3}p_1^0$	11.10	9009000 10.10	10.10	0001066	9.23	9901000 9.23 10834000
" $2s^2 2p^5 4d P_1^0$	10.97	9116000	6.97	10030000 9.11	9.11	10977000

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