

CALCULATION COVER SHEET

CALC NO. BOW-05-R9-15-03-00 DISCIPLINE Hydrology NO. OF SHEETS 5

PROJECT: UMTRA Site Characterization

SITE: Bowman, North Dakota

FEATURE: Volumetric Mixing Calculations: to estimate resultant concentrations of hazardous constituents in groundwater in lignite zone at point of compliance beneath Bowman disposal cell.

SOURCES OF DATA:

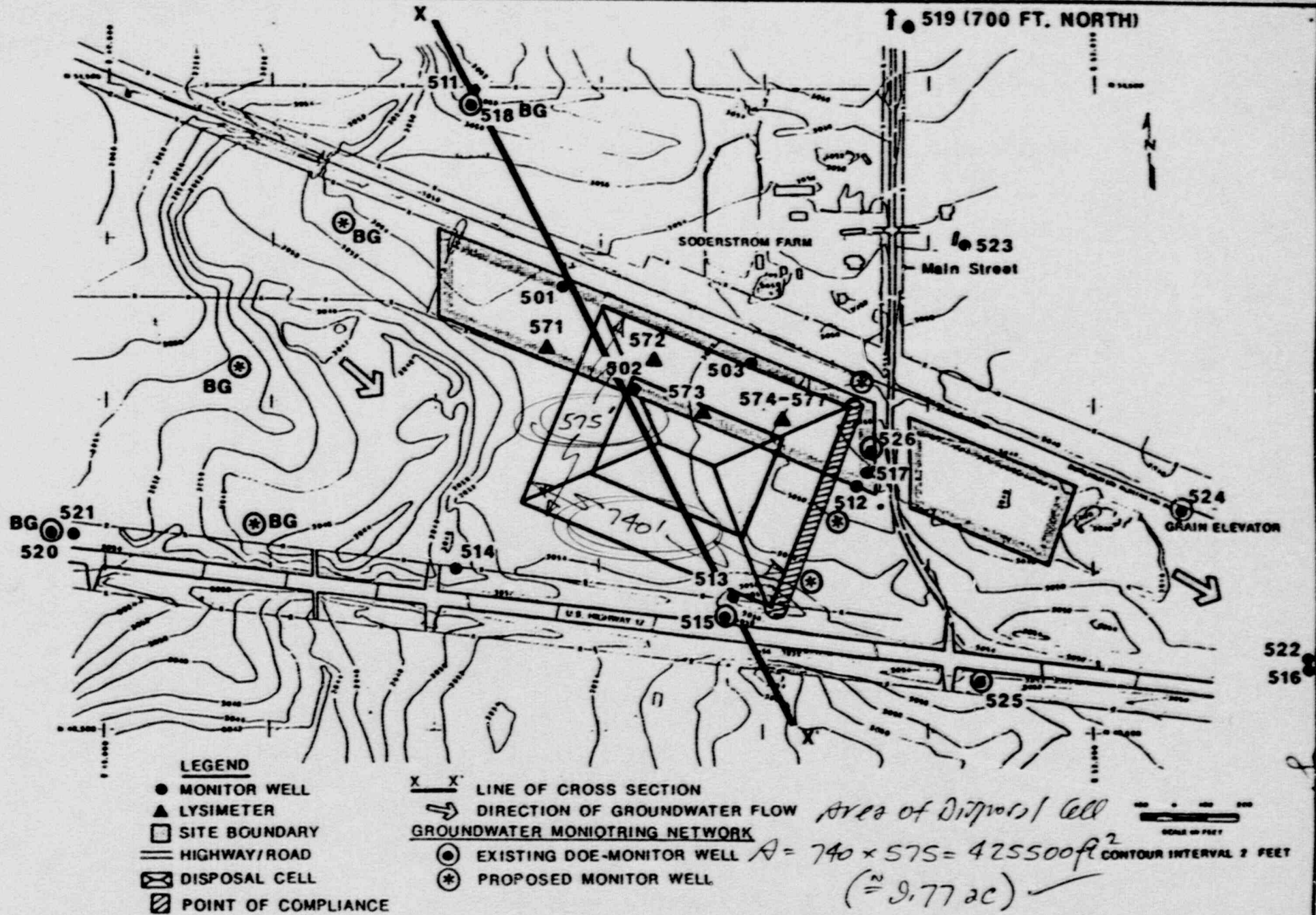
Source concentrations - from suction lysimeter water quality analyses.
Background water quality - from monitor well water quality analyses.
Cross-sectional area derived from figure E.2.1

SOURCES OF FORMULAE & REFERENCES:

Hem, J. D., 1985. Study and Interpretation of The Chemical Characteristics of Natural Water, USGS ~~Professional Paper~~ Water Supply Paper 2254.

PRELIMINARY CALC. FINAL CALC. SUPERSEDES CALC. NO. _____

		<u>R.J. Henderson</u>	<u>29 Aug 89</u>	<u>D. Matulis</u>	<u>9-15-89</u>		
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	<u>PDR WASTE</u>						
	<u>WM-74</u>	<u>PDC</u>					
REV. NO.	REVISION	CALCULATION BY	DATE	CHECKED BY	DATE	APPROVED BY	DATE



- LEGEND**
- MONITOR WELL
 - ▲ LYSIMETER
 - SITE BOUNDARY
 - == HIGHWAY/ROAD
 - ▨ DISPOSAL CELL
 - ▨ POINT OF COMPLIANCE

- X-X LINE OF CROSS SECTION
- ➔ DIRECTION OF GROUNDWATER FLOW
- GROUNDWATER MONITORING NETWORK**
- EXISTING DOE-MONITOR WELL
 - * PROPOSED MONITOR WELL

Area of Disposal Cell

$A = 740 \times 575 = 425500 \text{ ft}^2$

$(\approx 9.77 \text{ ac})$

SCALE IN FEET

CONTOUR INTERVAL 2 FEET

FIGURE E.2.1

**PLAN VIEW OF DISPOSAL CELL, POINT OF COMPLIANCE, AND GROUNDWATER MONITORING NETWORK
BOWMAN DISPOSAL SITE - GRIFFIN, NORTH DAKOTA**

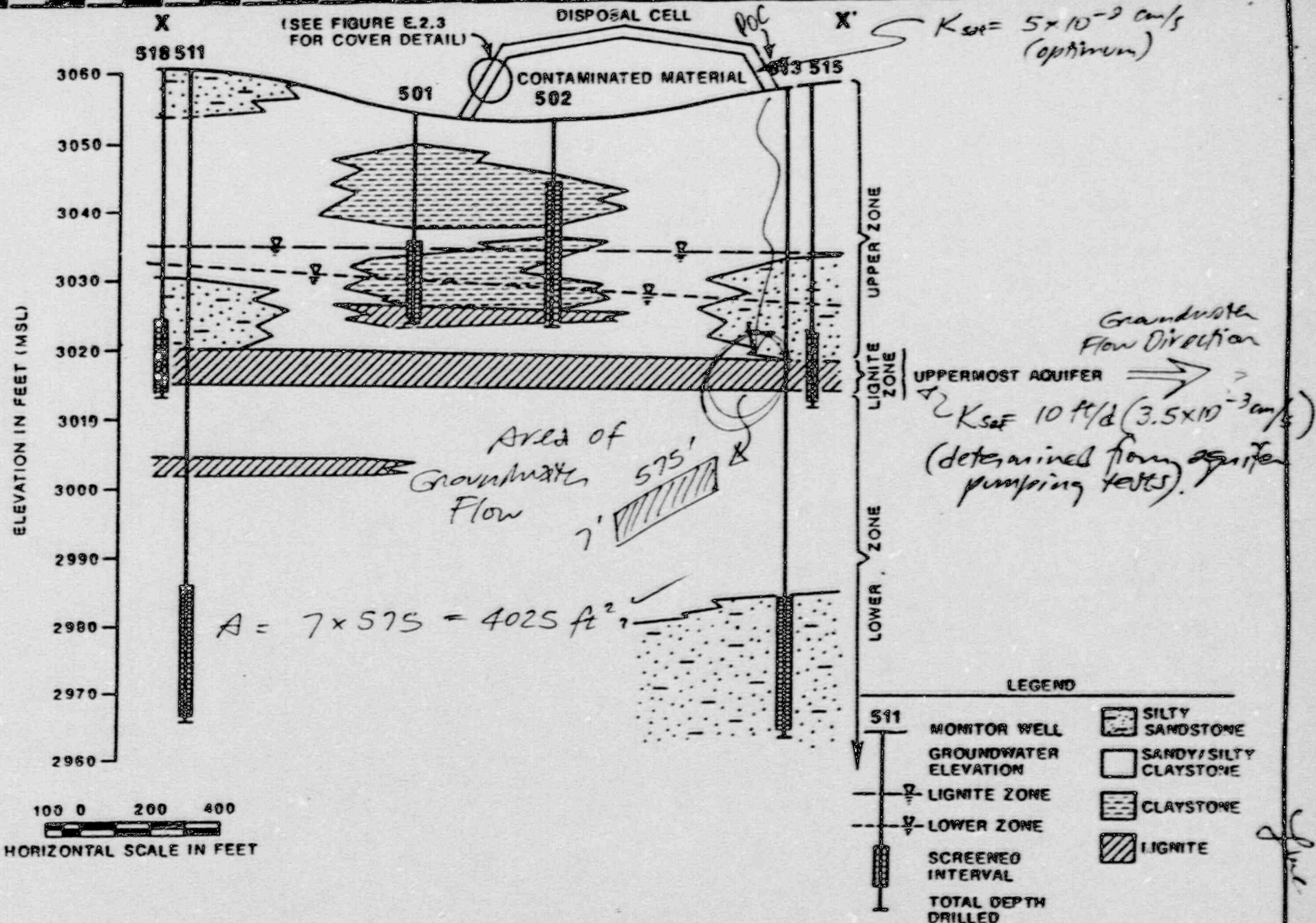


FIGURE E.2.2
 DIAGRAMMATIC CROSS SECTION OF PROPOSED DISPOSAL CELL
 BOWMAN DISPOSAL SITE - GRIFFIN, NORTH DAKOTA

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Line

Table E.3.2 Concentrations of designated hazardous constituents, Bowman disposal site - Griffin, North Dakota

Hazardous constituents ^a	MCL ^b (mg/l)	Proposed concentration limit ^a (mg/l)	<i>C_{GW}</i> Background-lignite zone ^c		<i>C_T</i> Source ^d concentration (mg/l)	<i>C_R</i> Groundwater concentration ^e (mg/l) hydraulic conductivity of cover (cm/s)	
			mean (mg/l)	range (mg/l)	5 x 10 ⁻⁹	1 x 10 ⁻⁸	
Antimony (Sb)	NS	0.02	0.02	0.09	0.09	0.02	0.03
Cadmium (Cd)	0.01	0.01	0.002	0.009	0.011	0.003	0.003
Chromium (Cr)	0.05	0.06	0.06	0.09	0.11	0.06	0.07
Lead (Pb)	0.05	0.05	0.01	0.05	0.05	--	--
Molybdenum (Mo)	0.10	0.10	0.05	0.07	0.20	0.06	0.07
Selenium (Se)	0.01	0.017	0.017	0.029	0.098	0.023	0.027
Uranium (U)	0.044	0.044	0.004	0.007	0.340	0.027	0.067
Vanadium (V)	NS	0.02	0.02	0.04	0.07	0.02	0.03

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^aSee Table E.3.1.

^bMCL: NS = not specified; MCLs from draft final EPA groundwater protection standards for UMTRA sites (40 CFR 192).

^cMean concentration from monitor wells 518 and 520.

^dMean concentration from lysimeters 572, 573, 575, 576, and 577.

^eBased on volumetric mixing calculations (Mem, 1985).

[Signature]

DATE 24 Aug 89

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BY RJH CHKD gmc

BOW-05-89-15-03-00

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A. Purpose of Calculation:

Perform volumetric mixing calculations to estimate resultant concentrations of hazardous constituents in groundwater in lignite zone at point of compliance beneath Bowman processing site.

B. Methods and Procedures:

Resultant concentrations were determined by using a solute-balance equation (Ham, 1985), which considers the volume of water infiltrating through the infiltration/leach barrier under a design saturated hydraulic conductivity of 1×10^{-8} cm/s (and an optimum of 5×10^{-9} cm/s), the source concentrations of hazardous constituents present in the stabilized material, the rate of discharge of groundwater through the lignite zone beneath the disposal cell, and mean background concentrations determined for hazardous constituents in the lignite zone.

C. Data Sources:

(Cr)
Source concentrations₁ - from suction lysimeters 572, 573, 575, 576 and 577, Bowman processing site. - water quality analyses.

(Cw)
Background water quality₁ from monitor wells 518 and 520 (lignite zone) - water quality analyses.

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D. Calculations and Analyses:

solute-balance equation:

$$C_R = \frac{(Q_{GW} \cdot C_{GW}) + (Q_T \cdot C_T)}{Q_{GW} + Q_T}$$

mg/L { C_R = resultant concentration in groundwater
 C_{GW} = background concentration in groundwater
 C_T = source concentration in contaminated materials

ft³/d { Q_{GW} = volume of groundwater flow in lignite zone
 Q_T = volume of water flow infiltrating infiltration/radon barrier

$$Q_{GW} = KiA$$

K = hydraulic conductivity = 10 ft/d
 i = hydraulic gradient = 0.002
 A = area = 4025 ft²

$$Q_{GW} = 10 \text{ ft/d} \cdot 0.002 \cdot 4025 \text{ ft}^2 = 80.5 \text{ ft}^3/\text{d}$$

$$Q_T = KiA$$

$K = 1 \times 10^{-8} \text{ cm/s} (2.8 \times 10^{-5} \text{ ft/d}), 5 \times 10^{-9} \text{ cm/s} (1.4 \times 10^{-5} \text{ ft/d})$
 $i = 1$ (unity) (assumption for steady state seepage) J_{se}
 $A = 425500 \text{ ft}^2$

$$Q_T(1 \times 10^{-8} \text{ cm/s}) = 2.8 \times 10^{-5} \text{ ft/d} \cdot 1 \cdot 425500 = 11.9 \text{ ft}^3/\text{d}$$

$$Q_T(5 \times 10^{-9} \text{ cm/s}) = 1.4 \times 10^{-5} \text{ ft/d} \cdot 1 \cdot 425500 = 6.0 \text{ ft}^3/\text{d}$$

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$C_{GW} + C_T = C_R$ values: see attached Table E.3.2

Calculations:

$$C_R = \frac{(Q_{GW} \cdot C_{GW}) + (Q_T \cdot C_T)}{Q_{GW} + Q_T} \quad (\text{in } \underline{\underline{mg/l}})$$

Chromium

Cd $1 \times 10^{-8} \text{ cm/s}$ $C_R = \frac{(80.5 \cdot 0.002) + (11.9 \cdot 0.011)}{80.5 + 11.9} = 0.003 \checkmark$

$5 \times 10^{-9} \text{ cm/s}$ $C_R = \frac{(80.5 \cdot 0.002) + (6.0 \cdot 0.011)}{80.5 + 6.0} = 0.003 \checkmark$

Chromium

Cr $1 \times 10^{-8} \text{ cm/s}$ $C_R = \frac{(80.5 \cdot 0.06) + (11.9 \cdot 0.11)}{80.5 + 11.9} = 0.07 \checkmark$

$5 \times 10^{-9} \text{ cm/s}$ $C_R = \frac{(80.5 \cdot 0.06) + (6.0 \cdot 0.11)}{80.5 + 6.0} = 0.06 \checkmark$

Molybdenum

Mo $1 \times 10^{-8} \text{ cm/s}$ $C_R = \frac{(80.5 \cdot 0.05) + (11.9 \cdot 0.20)}{80.5 + 11.9} = 0.07 \checkmark$

$5 \times 10^{-9} \text{ cm/s}$ $C_R = \frac{(80.5 \cdot 0.05) + (6.0 \cdot 0.20)}{80.5 + 6.0} = 0.06 \checkmark$

Selenium

Se $1 \times 10^{-8} \text{ cm/s}$ $C_R = \frac{(80.5 \cdot 0.017) + (11.9 \cdot 0.098)}{80.5 + 11.9} = 0.027 \checkmark$

$5 \times 10^{-9} \text{ cm/s}$ $C_R = \frac{(80.5 \cdot 0.017) + (6.0 \cdot 0.098)}{80.5 + 6.0} = 0.023 \checkmark$

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U_{uv} Ammonia
 $1 \times 10^{-8} \text{ cm/s} \quad CR = \frac{(80.5 \cdot 0.004) + (11.9 \cdot 0.340)}{80.5 + 11.9} = 0.047 \checkmark$

$5 \times 10^{-9} \text{ cm/s} \quad CR = \frac{(80.5 \cdot 0.004) + (6.0 \cdot 0.340)}{80.5 + 6.0} = 0.027 \checkmark$

5b Antimony
 $1 \times 10^{-8} \text{ cm/s} \quad CR = \frac{(80.5 \cdot 0.02) + (11.9 \cdot 0.09)}{80.5 + 11.9} = 0.03 \checkmark$

$5 \times 10^{-9} \text{ cm/s} \quad CR = \frac{(80.5 \cdot 0.02) + (6.0 \cdot 0.09)}{80.5 + 6.0} = 0.02 \checkmark$

Vanadium
 $1 \times 10^{-8} \text{ cm/s} \quad CR = \frac{(80.5 \cdot 0.02) + (11.9 \cdot 0.07)}{80.5 + 11.9} = 0.03 \checkmark$

$5 \times 10^{-9} \text{ cm/s} \quad CR = \frac{(80.5 \cdot 0.02) + (6.0 \cdot 0.07)}{80.5 + 6.0} = 0.02 \checkmark$

E. Results:

Results of the calculations are summarized in Table E.3.2 (attached).