# **Chapter 1-HW**

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ENGI-111-01

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11 February 2024

```
x = 10
y = 3
% a.
u = x+y
% b.
v = x*y
% C.
w = x/y
% d.
z = sin(x)
r = 8*sin(y)
s = 5*sin(2*y)
x =
    10
y =
     3
u =
    13
```

```
v =
     30

w =
     3.3333

z =
     -0.5440

r =
     1.1290

s =
     -1.3971
```

```
x = 5
y = 2
% a.
(1-1/x^5)^-1
% b.
3*pi*x^2
% C.
3*y/(4*x-8)
% d.
4*(y-5)/(3*x-6)
x =
     5
y =
     2
ans =
   1.0003
```

```
ans =
    235.6194

ans =
    0.5000

ans =
    -1.3333
```

```
a = 1.12
b = 2.34
c = 0.72
d = 0.81
f = 19.83
x = 1+(a/b)+(c/f^2)
s = (b-a)/(d-c)
r = 1/(1/a+1/b+1/c+1/d)
y = a*b*1/c*f^2/2
a =
  1.1200
b =
  2.3400
c =
  0.7200
d =
  0.8100
f =
  19.8300
```

```
x =
    1.4805

s =
    13.5556

r =
    0.2536

y =
    715.6766
```

```
a = 6.49
b = 0.0562
R = 0.08206
T = 273.2
V = 22.41
n = 1
P = n*R*T/V
P = n*R*T/(V-(n*b))-(a*n^2)/V^2
% The molecular attractions is the main cause of the difference in the two
% pressure estimates, as the term that is a correction for molecular
% attractions in the van der Waals equation, an^2/V^2, using the given
% values equals about equals about 0.013, while the difference between the
% first term of the van der Waals equation with the term that is a
% correction for the volume of the molecules, nb, and the ideal gas law is
% about 0.0025.
a =
    6.4900
b =
    0.0562
R =
```

```
0.0821

T = 273.2000

V = 22.4100

n = 1

P = 1.0004

P = 0.9900
```

```
a.

exp(-2.1^3)+3.47*log(14)+287^(1/4)
% b.
3.4^7*log(14)+287^(1/4)
% c.
cos(4.12*pi/6)^2
% d.
cos(4.12*pi/6)^2

ans =
13.2736

ans =
0.3062
```

```
ans = 0.3062
```

```
roots([60,20,-15,30])
ans =
    -1.0381 + 0.0000i
    0.3524 + 0.5979i
    0.3524 - 0.5979i
```

Published with MATLAB® R2023b