

CORRECTED COPY

Form C-103
(Revised 3-55)NEW MEXICO OIL CONSERVATION COMMISSION
MISCELLANEOUS REPORTS ON WELLS

(Submit to appropriate District Office as per Commission Rule 1106)

COMPANY Magnolia Petroleum Company, P. O. Box 2406, Hobbs, New Mexico
(Address)LEASE Lightcap Land Company WELL NO. 2 UNIT B 5 25 T 73 R 29E
DATE WORK PERFORMED 9-23-58 POOL UndesignatedThis is a Report of: (Check appropriate block) ☒ Results of Test of Casing Shut-off☒ Beginning Drilling Operations☐ Remedial Work☐ Plugging☐ Other _____

Detailed account of work done, nature and quantity of materials used and results obtained.

Commenced drilling operations 10:30 a.m. September 22, 1958 (spud date).

Set 413' of 13-3/8" csg. on bottom, cemented w/400 sks. neat. Circulated out 45 sks.
Plug down 12:30 p.m. September 23, 1958. WOC 24 hours. Tested 13-3/8" csg. w/500#
before and 300# after drilling plug, held O.K.

FILL IN BELOW FOR REMEDIAL WORK REPORTS ONLY

Original Well Data:

DF Elev. _____ TD _____ PBD _____ Prod. Int. _____ Compl Date _____

Tbng. Dia _____ Tbng Depth _____ Oil String Dia _____ Oil String Depth _____

Perf Interval (s) _____

Open Hole Interval _____ Producing Formation (s) _____

RESULTS OF WORKOVER:

BEFORE

AFTER

Date of Test

Oil Production, bbls. per day

Gas Production, Mcf per day

Water Production, bbls. per day

Gas-Oil Ratio, cu. ft. per bbl.

Gas Well Potential, Mcf per day

Witnessed by _____

(Company)

OIL CONSERVATION COMMISSION

I hereby certify that the information given
above is true and complete to the best of
my knowledge.Name M. L. ArmstrongName Myung, Jr.

Title _____

Position District SuperintendentDate Oct 1 1958Company Magnolia Petroleum Company

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Opinion: _____

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1. *Chlorophyll a* (Chl *a*)

Figure 1. Schematic representation of the experimental design. The subjects were divided into two groups: the control group (n = 10) and the experimental group (n = 10). The control group received a placebo (P) and the experimental group received a 10% solution of the active ingredient (A). The subjects were divided into two groups: the control group (n = 10) and the experimental group (n = 10). The control group received a placebo (P) and the experimental group received a 10% solution of the active ingredient (A). The subjects were divided into two groups: the control group (n = 10) and the experimental group (n = 10). The control group received a placebo (P) and the experimental group received a 10% solution of the active ingredient (A).

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