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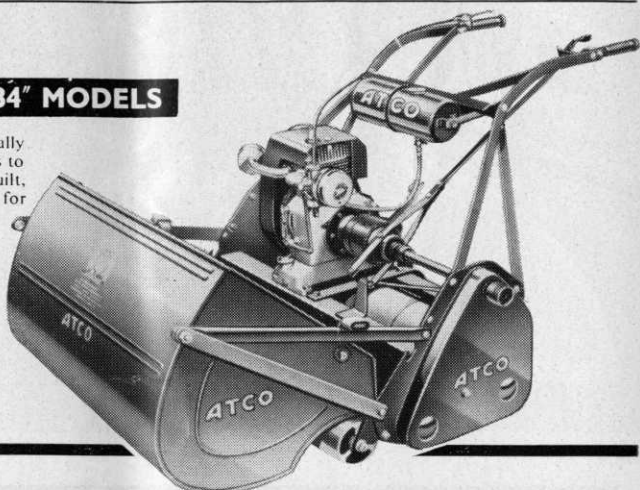
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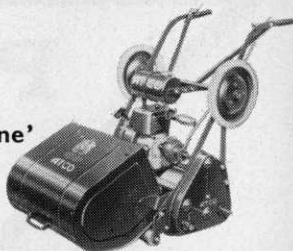
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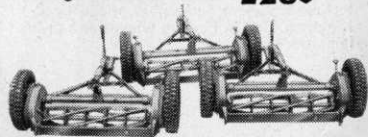
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No. 245 New Series

AUGUST 1965

REASONING

*The air is blue around the tee—
Invectives rise and fall;
Our shortest-tempered member must
Have just addressed the ball.*

—A. R. FONTENOT

AUGUST

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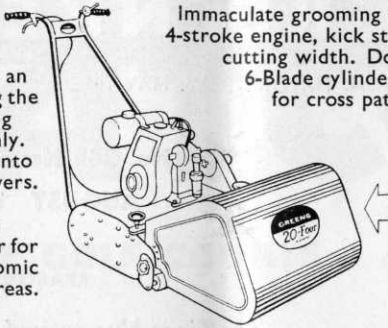
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TEE SHOTS



by the Editor

Progress

The leading bodies in British golf set up a Golf Development Council at the C.C.P.R. offices in London on 22nd July. Stated objects include co-operation with government departments and other bodies to stimulate the provision of more golf facilities and encourage teaching the game in schools.

Mr. Gerald Micklem was elected Chairman and his wide experience of the game will be invaluable in consolidating and developing this important advance. The R. & A., national golf Unions, Golf Foundation, P.G.A., L.G.U., Public Courses Association and Artisan Golfers Association were all concerned in the inaugural meeting and are contributing to the funds.

* * *

More Bounce to the Ounce

British golf ball manufacturers are said to have surprised the Americans with the distance some of their new balls will travel. There is no initial velocity test in Britain. 250 feet per second is the maximum allowed by the U.S.G.A. Distance, however, still has to take its place with consistent flight, feel, and sound and, of course, durability.

* * *

Moving

Mr Bill Payne has moved from Sutton Coldfield to the south-west and will be living in North Street, Cheddar. Warwickshire greenkeepers will be glad to know that their county is still in his territory.

* * *

Careful

E. Hanlon, who lives at Avon in Connecticut, holed his tee shot on the 170 yard 11th at the Avon Country Club. He declared his ball in the hole "unplayable", picked it out of the cup, and putted in for a three. Amateurs in the U.S. are allowed to take \$200 worth of merchandise from firms which hand out gifts for "aces", but Hanlon would rather see the money go to some kind of charity than over the bar to entertain admirers.

A

PERENNIAL

PROBLEM

by **H. J. LIDGATE**

*Chemist, The Sports Turf Research
Institute*

IN the months of August and September we can usually expect to receive a few inches of rain which will begin to moisten the soil after the fairly dry summer. From many points of view this rain is most desirable but there are inevitably snags in all good things. The greenkeeper will find that in addition to a greater amount of grass cuttings in the mower's box he may well have an extra helping of worm casts also.

Casting

The earthworm is brought up by autumn rain from the deeper regions of the soil where it spends the dry summer and once again begins to cast on the surface. This is most unfortunate when the soil surface in question happens to be one of your prize golf greens or on a fairway or tee. It is a great pity that some, though not all, earthworms have this bad habit of casting on the surface because they do help a great deal in aerating the soil with their burrows. These burrows also help the penetration of moisture into the soil. At St Ives there is a difference between the drought resistance of plots treated with wormkiller (lead arsenate) and untreated plots, the latter suffering a lot less from drought in dry weather. However, the troubles caused by allowing the earthworms to work unchecked more than counter any possible benefits.

On greens the activities of the earthworm are most unsightly and result in an increase of annual meadow grass and weeds in the turf. On tees the resulting muddiness of the surface leads to heavier wear and the need for more renovation whilst on fairways the effect of large numbers of casts is to give a greasy surface which is unpleasant for walking on and can make slopes very slippery.

At most golf courses some wormkilling measures are taken though they are sometimes of limited extent. The usual difficulty is in the expensiveness of the presently available control measures. Lead arsenate, that old standby of so many greenkeepers now costs approximately £200 per ton, i.e. about £5 to £7 per green according to the size of the green and approximately £60 per acre for the treatment of fairways excluding the cost of labour involved in applying the material. Of course, this should give good results for several years — maybe six or more — but we do hear reports of the material not always working in which case it is an expensive failure. It is safer to carry out a trial of say $\frac{1}{2}$ cwt. and observe results in twelve months time than to go ahead with larger quantities in the first place. Calcium arsenate which is a rather cheaper material is now unfortunately unobtainable as nobody manufactures it any longer.

Choice

There are alternative materials, Chlordane for instance is very popular at present though again we get odd reports of complete failure even when the right amount of Chlordane has been used. It costs approximately £12 per acre for the material, i.e. £1 to £1 10s. per green and it usually gives good results for at least one year. For those who prefer to use a less poisonous material, Derris dust or Mowrah meal are available each costing about £27 per acre excluding the cost of the labour involved — rather a lot with Mowrah meal which requires watering in with a high pressure hose to get the best results,

followed by sweeping up of the dead worms afterwards. The results obtained from these materials depend very much on the timing of the work. It is important that the material shall be used when the worms are very near the surface preferably in warm moist weather as often occurs in October. The treated areas may be free from worms for up to two years though one year is more usual. Neither of these products persist in the soil and where only small areas are treated worms soon re-invade from the surrounding areas. Where a good initial kill of worms is achieved over a large area then it may remain worm free for a year or two until the worm population builds up again.

Persistent

Copper sulphate has been used occasionally in the past as a worm-killer with success but there is some risk of damaging the turf if too much is applied. On areas of undulating turf there is a considerable risk of the copper sulphate running into hollows and causing the grass to be killed in these low parts. The copper remains in the soil and the subsequent re-establish-

ment of grass is difficult. As a result of this one hesitates to recommend the use of copper sulphate for wormkilling.

Caution

Irrespective of which material is used the greenkeeper will have to do the actual job of applying it. It is up to him to see that the right amount of material is used. It is not reasonable to expect an amount of material sufficient for one green to treat three or four greens and still give good results. It is very helpful if a careful note is kept of the quantities applied to each area and if possible a small sample of the material used should be kept. If the results are not all they should be it will be possible to check how much worm-killer has been used and also to check the quality of the material used. Finally, the greenkeeper should always take great care when handling these materials especially the more poisonous ones such as lead arsenate, calcium arsenate or chlordane, taking care to avoid working in dust or spray drift and to wash thoroughly before eating or smoking and to wash in any case when the job has been finished.

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Continuing . . .

WHY TURFGRASS RESPONDS TO PROPER USE OF LIME AND SULFUR COMPOUNDS

By Dr **ELIOT C. ROBERTS**
Turfgrass Specialist,
Iowa State University,
Ames, Iowa

*With grateful acknowledgements to
"The Golf Course Reporter"*

Phosphorus

Soil pH has a great effect on the availability of phosphorus to plants. Phosphates are most available from pH 5.5 to 7.5. Above and below these levels phosphates are tied up with other minerals and their availability is reduced. Under acid conditions iron and aluminium phosphates are formed. The solubility of iron and aluminum compounds increases as soils become more acid until they are readily soluble at pH levels from 4.5 to 5.0. Under these conditions the iron and aluminum react with phosphorus and convert it to a form which plants cannot use.

At pH levels above 7.5 phosphates again become less available. It is believed that under these conditions phosphorus is tied up as insoluble calcium phosphates which are unavailable for plant use. In the pH range 5.5 to 7.5 calcium phosphates of a type which are more mobile and available for plant use are formed. Within this range it is likely that the ability of the root to absorb sufficient phosphorus is more responsible for nutritional status of the plant than the availability of soil and fertilizer phosphorus.

It should be noted that at very low soil pH levels increased solubility of aluminum, iron, copper and manganese may result in higher concentrations of these elements than can be tied up with phosphorus. In such instances toxicity to plants may develop. Aluminum toxicity may become particularly serious as excessive amounts of this element are absorbed and precipitated in the conducting tissue as aluminum hydroxide. Deficiencies of copper, zinc, iron and manganese are more likely to occur under alkaline soil conditions.

Potassium

The availability of potassium does not change to any great extent over the normal soil pH range. There is some indication that heavy liming has reduced availability slightly and that under acid soil conditions potassium leaches from the root zone more readily.

Calcium and Magnesium

Calcium availability increases as soils become less acid and more alkaline. In general the quantity of calcium available in the average soil at pH 4, 5, 6, 7, and 8 is relatively as 1 : 4 : 8 : 14 : 22 respectively.

Magnesium availability increases as soils become less acid and more alkaline up to pH 8.5 after which it decreases. In general the quantity of magnesium available in the average soil at pH 4.5, 5.5, 6.5, 7.5, and 8.5 is relatively as 1 : 2 : 3 : 4 : 2½ respectively. The rate of increase for magnesium over the pH range of soil is therefore much smaller than for calcium.

Under acid soil conditions calcium and magnesium may be deficient and as such contribute to unsatisfactory plant growth.

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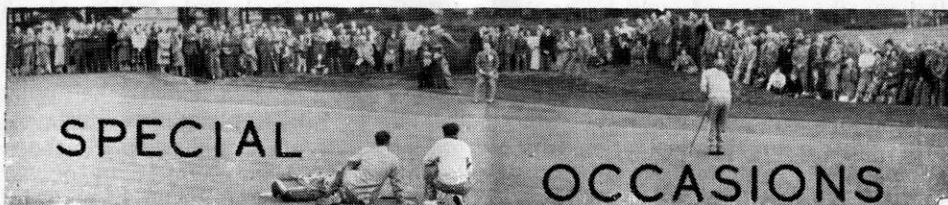
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29th	East Midland Section, Autumn Tournament, Birstall Golf Club.
29th	Welsh Section, Autumn Tournament, Neath Golf Club.
OCTOBER 7th	S.G.G.A. West Section, Autumn Meeting, Pollock.
19th	Northern Section, Autumn Tournament, Leeds Golf Club.
29th	Midland Section, Annual Dinner, King's Head Hotel, Bearwood.

Manganese and Iron

Besides soil pH, soil air and water content, degree of soil compactness and soil organic matter are related to availability of iron and manganese. Since plants require only small amounts of these two nutrients for satisfactory growth, toxicity from their over abundance may be as great a problem as deficiency from their short supply. Excess supply is likely to occur under extremely acid conditions while deficiencies are usually noted within the pH range 7.5 to 8.5. At high pH levels iron deficiency may occur because of an unfavourable iron-calcium balance within the plant which renders the absorbed iron ineffective. In this instance sufficient iron is absorbed, but utilization is prevented.

These deficiencies occur more on sandy soils than on heavier soils and are more likely to be noted in spring and early summer than later in the year.

How to make Soils less Acid

When soil pH levels drop below 5.8, treatments should be made to elevate the pH to 6.5-7.5. Lime is the material most often used for this purpose. Calcitic limestone contains calcium car-

bonates. Dolomitic limestone contains calcium carbonates and magnesium carbonates (equivalent to 15 to 20 percent magnesium oxide). Limestone may be broken down into calcium oxide and carbon dioxide. Calcium oxide is called quicklime. Quicklime plus water yields hydrated lime. Hydrated lime and quicklime are more soluble and faster acting than ground limestone. Mixtures of finely ground limestone and hydrated lime are called Agricultural lime. Actually soil acids in the presence of limestone or hydrated lime produce calcium and magnesium ions which are attracted to the clay particles in exchange for hydrogen ions which react with the carbonate to form carbonic acid. Carbonic acid is weak and slowly breaks up to form carbon dioxide and water.

The chemical composition of lime determines its neutralizing power. The relative neutralizing power of lime has been calculated on the basis of pure calcium carbonate being equal to 100 percent. Accordingly

- 100 lb. of calcium carbonate
- 74 lb. of calcium hydroxide
- 88 lb. of magnesium carbonate
- 56 lb. of calcium oxide
- 40 lb. of magnesium oxide —

all have the same value in correcting soil acidity. The following tables give percent neutralizing value and lb. equivalent to one ton of calcium carbonate.

In addition to chemical composition the particle size of lime affects its reaction in the soil. The finer the particles the faster they react; the coarser the particles the slower they react. A good lime should have some fine particles for immediate reaction and some coarse particles to add residual effect to a treatment. In general a reasonably fine limestone will have a

it is to change it abruptly with one large application. In general the finer the grind the lighter the application or the coarser the grind the heavier the application. Also, applications of finely ground material should be made more often because lasting effects are not as great. Very coarse material may be used on fairways at rates from 150 to 200 lb. per 1,000 square feet. On the basis of fine ground limestone, a safe rule to follow is — increase the rate of application 25 to 50 percent for coarse materials and reduce it 25 to 30 percent for hydrated materials.

Material	Molecular Weight	Neutralizing Value — %	Lb. Equivalent to 1 ton Calcium carbonate
CaCO ₃ Calcium Carbonate	100	100	2,000
MgCO ₃ Magnesium Carbonate*	84	119	1,680
Ca(OH) ₂ Calcium Hydroxide	74	135	1,480
Mg(OH) ₂ Magnesium Hydroxide	58	172	1,160
CaO Calcium Oxide	56	178	1,120
MgO Magnesium Oxide	40	250	800

particle size analysis similar to the following:

100 percent through a 10 mesh¹ screen
75 percent through a 50 mesh screen
60 percent a 100 mesh screen

In using these materials it is well to remember that hydrated lime is fine and thus more soluble in water and faster acting. It is more likely to burn and should never be applied on fairways at rates larger than 50 lb./1,000 square feet. Heavy applications at one time cause the soil at the surface (where the majority of turfgrass roots are located) to become too alkaline. This may tie up phosphorus, iron and other minor elements until the reaction works down into the soil. It is far better to make several smaller applications of lime (25 to 50 lb./1,000 square feet) and change the soil pH gradually than

Late fall, winter, and early spring are the best times to lime. Treatments made while the soil is frozen in the winter usually work well. Spreading lime at this time keeps additional spreader traffic off the turf during the growth season. Never use hydrated lime on fairways at more than 25 lb. per 1,000 square feet during the summer. If necessary other limes may be applied at standard rates during the summer. Never use hydrated lime before or after the application of a fertilizer containing ammonia. The lime will activate the fertilizer and produce ammonia gas which is extremely toxic to turfgrass. It is desirable to lime well in advance of making applications of phosphorus and arsenic. Where treatments with these materials follow closely, an application of lime, they will be less effective because of increased fixation in the soil.

Putting greens should be limed with

¹ Ten openings per linear inch; 100 openings per square inch.

fine ground limestone at from 10 to 50 lb. per 1,000 square feet. Hydrated lime should not be used on greens at rates in excess of 20 lb./1,000 square feet. Where greens have become scalded during the summer, anaerobic soil conditions may be cleaned up with applications of 2 to 5 lb. of hydrated lime per 1,000 square feet. Treatments may be made every week until a total of 10 lb./1,000 square feet has been applied. The material may be mixed with sand to increase ease of spreading.

In addition to lime, gypsum (calcium sulfate) may be used to supply calcium to acid soils. The effect of gypsum on soil pH is one of initial aciduation followed by a tendency for the pH to rise. Soils with poor structure which are slow to drain have been improved by applications of gypsum. Seventy-five to 100 lb. of gypsum per 1,000 square feet worked into wet spots in fairways have helped correct some poor drainage areas.

What determines how much Lime is needed

Since the concentration of hydrogen ions in the soil is responsible for its acidity it is necessary to consider both those ions attached to soil particles as well as those in soil solution. Hydrogen ions in solution constitute the active acidity of a soil. Hydrogen ions attached to soil particles make up the reserve acidity. If only sufficient lime is added to neutralize the soil solution, hydrogen ions will quickly be exchanged from the soil particles to the solution and the pH will return to an acid reading. Only where sufficient lime is applied to neut-

ralize both the hydrogen ions in the solution and on the soil particles will a significant change in soil pH be brought about.

A measurement of soil pH gives only an indication of active acidity. It is necessary to run a lime requirement test in order to accurately evaluate the total acidity (active plus reserve acidity) so that the proper amount of lime may be applied. Lime requirement tests show that a sand and a clay soil may have the same pH but the clay soil requires more lime to bring about a given change in soil pH.

Thus heavier soils and soils with higher percentages of organic matter have higher lime requirements. As soils increase in the percentage of small particles contained (for example sandy soils to loamy soils to clay soils) increasing amounts of lime are required to counteract the reserve acidity. As soils increase in the amount of organic matter, they contain (for example sandy soils of low organic matter to sandy soils of medium organic matter) the quantity of lime required to neutralize the reserve acidity becomes greater. Actually the silt and clay and humus fractions on a soil serve as a buffer system which causes these soils to resist change in pH. Once limed to the proper pH level, they will be more stable and become acid more slowly than lighter textured soil of low organic matter content.

Finally, the more acid the soil the greater amount of lime required to raise the pH to a desired level. These relationships may be noted in Table 2.

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250	5	0.40
500	10	0.70
1,000	20	1.20

How to make Soils more Acid

Soils which have high pH values can be made more acid by the application of fertilizers containing ammonium nitrogen. This can be accomplished over a period of time on tees and greens by applying from $\frac{1}{4}$ to $\frac{3}{4}$ lb. nitrogen per 1,000 square feet per application. Frequent treatments are usually necessary to provide adequate nitrogen levels over the entire growth season. Attempts to increase the amount applied at each treatment so that the frequency of applications may be reduced often result in foliar burn or in the production of over succulent turf which is subject to injury from adverse climatic conditions. Applications of fertilizers containing ammonium nitrogen may be applied dry with a whirl-wind type spreader or may be spread with a proportioner using water as a carrier. In either case the fertilizer should be washed from the foliage and into the soil immediately following application to each green or tee.

Aluminum sulphate has been used to acidulate the soil; however, rates of application for turf have not been well worked out. This material can be extremely toxic to plants and its use on turf is not generally recommended.

Flowers of sulfur or powdered sulfur may be used effectively to acidulate the soil. It has been used on putting greens and tees where pH levels have been too high for best growth of turfgrasses. Most uniform application may be made by mixing the sulfur with sand or topsoil as a carrier and by spreading the mixture with a whirl-wind type applicator.

It is important that exactly the right amount be applied since sulfur decomposes to form sulfuric acid which in large amounts may produce severe injury to turfgrass stands. Tests with a very fine sandy loam soil to which powdered sulfur was mixed at rates from 100 lb./A to 1,000 lb./A produced the following results:

It was found that the maximum effect of the sulfur on reducing soil pH was evident within one month from the date of application. Tests with heavier soils and with soils higher in organic matter indicate less effect than on the lighter soils with lower organic matter contents.

Applications on turf should not exceed 5 lb./1,000 square feet. Treatments should be made in spring and fall rather than during summer months. Test strips on a turf nursery are advised prior to treating greens. These will enable a careful standardisation of rate of application with existing grass and growth conditions.

Since disease organisms are often more active under acid soil conditions, particular attention should be directed toward disease identification and fungus control during periods of soil acidulation with powdered sulfur.

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PREDICTION

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St Peter meets the Coach,
He'll hear "Now Peter, get this straight—
It's all in the approach"!*

—Anita Raskin