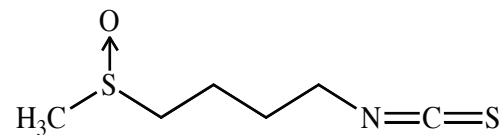
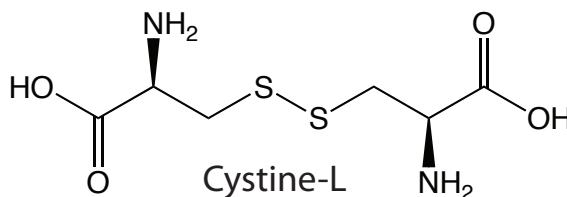


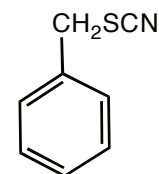
Isothiocyanates



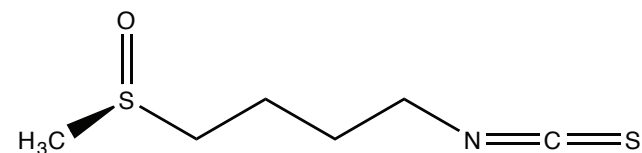
R, S-Sulforaphane



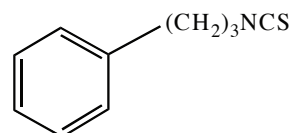
Cystine-L



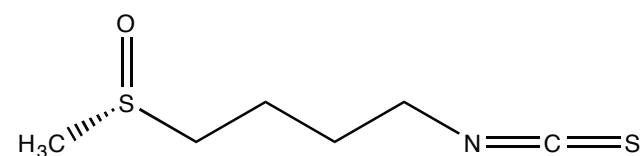
Benzyl Isothiocyanate



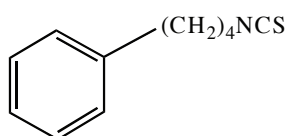
R-Sulforaphane



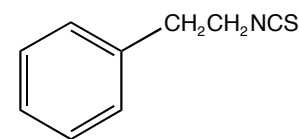
3-Phenylpropyl Isothiocyanate



S-Sulforaphane



4-Phenylbutyl isothiocyanate



Phenethyl Isothiocyanate

Isothiocyanates occur naturally as glucosinolates in edible plants.^{1,2} The consumption of cruciferous vegetables is the main dietary source of isothiocyanates.³ Benzyl and phenethyl isothiocyanates (BITC, PEITC) have been found to inhibit chemically induced carcinogenesis in several animal models.^{4,5} BITC is particularly effective against benzo[a]pyrene-induced lung tumorigenesis, while PEITC inhibits NNK-induced lung carcinogenesis.⁶ 3-Phenylpropyl and 4-phenylbutyl isothiocyanates (PPITC, PBITC) are synthetic compounds that show anti-carcinogenic effects.⁷ Studies on the effect of alkyl chain length of phenylalkyl isothiocyanates on tobacco specific nitrosamine-induced lung tumorigenesis revealed that PPITC and PBITC are more effective than the naturally occurring isothiocyanates.⁸

Sulforaphane, isolated from broccoli⁹, has been found to inhibit chemically induced mammary tumors in rats.¹⁰ Many sulforaphane analogues have been previously isolated from plants.¹¹ Their enzyme inducing activity was less potent than that of sulforaphane.⁹ Isothiocyanates have been found to induce activity of the detoxifying enzymes system, glutathione S-transferase^{5,12} and to suppress carcinogen activation by cytochromes P450^{4,5,13}, particularly P450 2B1, the major enzyme involved in NNK activation.¹⁴ LKT Laboratories, Inc. offers the natural chiral R-sulforaphane, its synthetic enantiomer, S-sulforaphane, and synthetic R,S-sulforaphane. Phenylalkyl isothiocyanates and phenethyl glucosinolate are also available in high purity.

1. Tookey, H. L.; et. al., Glucosinolates. In Toxic Constituents of Plant Food Stuffs, pp. 103-142 (1980).

2. Kjaer, A. Chemistry of Organic Sulfur Compounds, Vol. 1, pp. 409-420 (1961).

3. Fenwick, G. R.; Heaney, R. K.; Mullin, W. J. CRC Crit. Rev. Food Sci. Nutr. 18, 123-201 (1983).

4. Wattenberg, L. W. Carcinogenesis. 8, 1971-1973 (1987).

5. Chung, F.-L.; Jachatz, A.; Vitarius, J.; Hecht, S.S. Cancer Res. 44, 2924-2928 (1984).

6. Lin, J.M., Amin, S., Trushin, N., and Hecht, S.S. Cancer Lett. 74:151-159 (1993).

7. Zhang, Y., and Talalay, P. Cancer Res. 54(7 Suppl):1976S-1981S (1994).

8. Morse, M. A.; Eklind, K. I.; Amin, S. G.; Hecht, S. S.; Chung, F.-L. Carcinogenesis. 10, 1757-1759 (1989).

9. Zhang, Y., Talalay, P., Cho, C., and Posner, G.H., Proc. Natl. Acad. Sci. USA 89, 2399-2403 (1992).

10. Zhang, Y., Kensler, T.W., et al., Proc. Natl. Acad. Sci. USA 91, 3147-3150 (1994).

11. Kjaer, A., Fortschr. Chem. Org. Naturst. 18, 122-176 (1960).

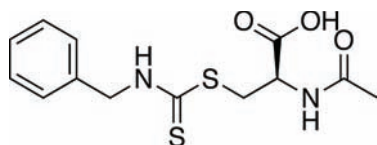
12. Benson, A. M.; Barretto, P. B. Cancer Res. 45, 4219-4223 (1985).

13. Morse, M. A.; Amin, S. G.; Hecht, S. S.; Chung, F.-L. Cancer Res. 49, 2894-2897 (1989).

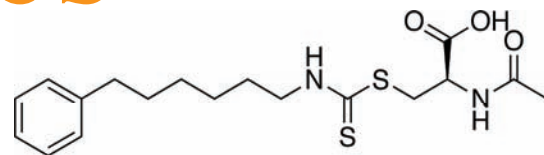
14. Conaway, C.C., Jiao, D., and Chung, F.L. Carcinogenesis, 17:2423-2427 (1996).



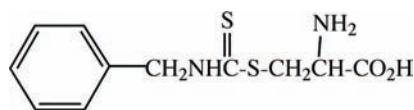
Isothiocyanate Conjugates



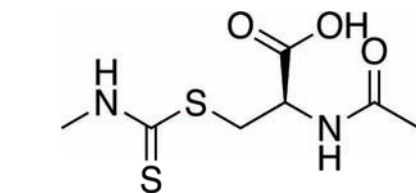
N-Acetyl-S-(N'-benzylthiocarbamoyl)-L-Cysteine



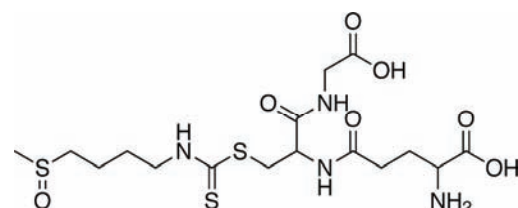
N-Acetyl-S-(N'-phenylhexylthiocarbomoyl)-L-cysteine



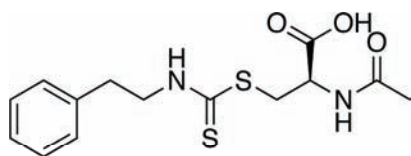
S-(N-Benzylthiocarbamoyl)-L-cysteine



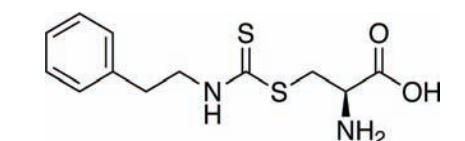
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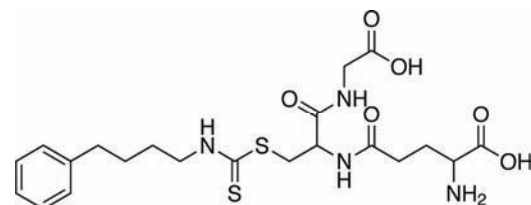
S-(N-Methylsulfinylbutylthiocarbamoyl)-glutathione



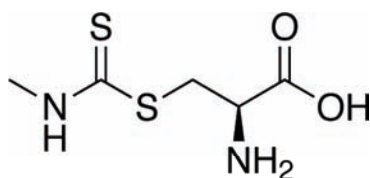
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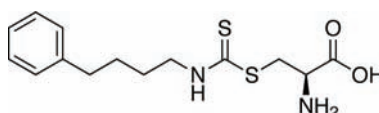
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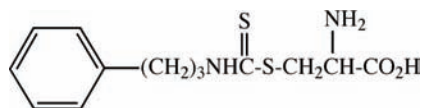
S-(N-Phenylbutylthiocarbamoyl)-glutathione



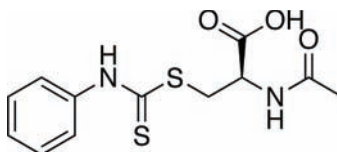
S-(N-Methylthiocarbamoyl)-L-cysteine



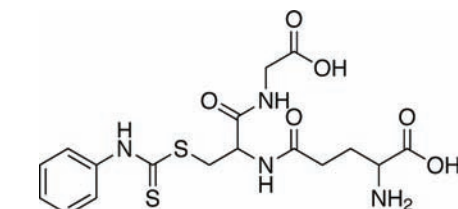
S-(N-Phenylbutylthiocarbamoyl)-L-cysteine



S-(N-3-Phenylpropylthiocarbamoyl)-L-cysteine



N-Acetyl-S-(N'-phenethylthiocarbamoyl)-L-cysteine



S-(N-Phenylthiocarbamoyl)-glutathione

