

Mechanical -strength characteristics of concrete made with stainless steel industry wastes as binders

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Abstract

One of the problems of steelworks, referring to the steel manufacture, is the generation of secondary products, resulting in recent years in new types of industrial waste that, depending on the sustainability of the environment, must be reused or treated. This paper analyzes the mechanical behaviour of concrete samples with the addition of an industrial waste, such as ferritic fume dust produced by electric arc furnaces (EAF) when the materials are melted and makes a comparison using other types of additions of concrete such us silica fume. At the same time this paper studies the capacity of the matrix to encapsulate this residue that eventually ends up deposited in a landfill. The results show that, besides giving the concrete a greater resistance as it happens with silica fume, the use of this type of waste as an addition to concrete is suitable since the material remains encapsulated in the concrete matrix, thus not producing leaching of heavy metals which can be harmful to the environment and therefore to the health of the human being.

Keywords: Concrete, Fume dust EAF, Compressive strength, Flexural strength

1. Introduction

One of the biggest problems today in the steel industry is the production of waste in varying amounts depending on the type of scrap used and steel to be processed, generating various types of steel including ferritic steel. Ferritic steel is a type of steel that is composed of less than 0.10% carbon. This grade of steel was developed as a stainless steel group (ACERINOX) that can resist oxidation and corrosion, specifically stress cracking corrosion (SCC). We will focus on the study of ferritic steel for our study. These materials produce large amounts of waste that build up fume dust. Said dust is generated in the combustion of the scrap inside the electric arc furnace (EAF). The powder evaporates and is extracted, then deposited in baghouse dust collectors for its later removal and deposition into landfill. These are expensive to recycle or reuse and difficult to transform into an inert waste.

Current steel production is about 1597 million tonnes according to International Iron and Steel Institute (IISI) Conferences 2015 [1]. EAF fume dust is generated in appreciable quantities when casting steel.

37 This transformation creates environmental problems in the steel industry. Today the reuse of waste in
38 the metallurgical industry does not exist or is inappropriate.

39 From an environmental point of view, the American Environment Agency (EPA) [2,3] classified these
40 materials as toxic and dangerous products, just as they are under Spanish legislation. The toxicity of
41 these materials is based on their content of non-ferrous metals, mainly, chromium, zinc, lead, nickel
42 and magnesium, all of them easily leachable metals with consequently harm caused to soil and
43 groundwater when stored in landfills.

44 The high content of these components requires us to seek solutions aside from treatment and/or
45 hazardous landfill. Furthermore, the economic importance of the recovery of certain metals (zinc...)
46 would be significant since the waste generated in the steel manufacture is composed of a large amount
47 of metals that can be recovered through certain processes that sometimes are cheaper than the
48 acquisition as raw material. Recovery rates of around 2% of the gross production have been verified
49 in the steel industry [4,5].

50 One solution is to incorporate the steelmaking waste in concrete, either completely or partially
51 recovered and after conducting the appropriate tests, verify the correct behaviour in the different
52 stresses to which the new material will be exposed. Different proportions of addition were studied.

53 Given its characteristics fume dust has binding properties if incorporated into appropriate systems so
54 that its reuse does not contaminate and exert a favourable effect on the system.

55 The results of the trials demonstrate out below improving the mechanical strength of the material.

56 One of the problems that industrial wastes can pose is when the solid materials come into contact with
57 a liquid,[2,6,7] some of its components can dissolve to a lesser or greater extent, being of interest the
58 degree of dissolution for each individual constituent. The leaching of materials can occur in the place
59 where it is applied, by natural infiltration of water, rain, exposure to seawater, etc.

60 The problem of environmental pollution can be established in several branches, waste water and waste
61 management are usually solved by solidification-stabilization [8–10].

62 Another of the materials used as addition and stabilization of materials is the slag electric arc furnace
63 in which the material acquires great importance in the recovery of waste [11–15].

64 There are several solidification-stabilization techniques, such as: the absorption of materials, which
65 commonly coagulate and precipitate to subsequently pass the solid phases or use cement to generate
66 concrete blocks containing this organic waste. There are authors who have used this procedure with
67 toxic waste from metals[16,17], or waste from the steel industry Norma UNE 83491 EX.

68 This stabilization presents great advantages for the treatment of hazardous waste with metals such as:
69 a low technical requirement in personnel, low production cost, long-term structural stability of the
70 concrete, high resistance to biodegradation, low permeability of concrete.

71 There are a large number of factors that influence the relationship in which the constituents of a
72 material are dissolved from the matrix material. These factors can be divided into physical and
73 chemical.

74 Currently there are several legislations in the European regulations in which are found IV.39/9 34
75 2000/532/CE, by which a list of hazardous waste is established.

76 According to Spanish regulations, those listed in Annex 2 of Royal Decree 952/1997, of June 20, where
77 the list of hazardous waste approved by Decision 94/904 / EC, of the council, of 22 December, in
78 accordance with section 4 of article 1 of directive 91/689 / EEC

79

80 The treatments can be classified as:

- 81 • Stabilization: Treatment of the waste with chemical agents, such as lime or phosphoric
82 acid, to reduce the mobility of the components.
- 83 • Solidification: Confine in a matrix of low permeability. This improves the mechanical
84 properties and decreases the contact surface of the waste with the possible leachant.
- 85 • Chemical extraction: Treatment that involves washing the metals contained in the ashes
86 with different liquids.

87

88 This type of treatment is focused on the immobilization of metals, since organic molecules are not
89 easily incorporated into crystalline structures and it is more difficult for them to form insoluble
90 precipitates. Stabilization and solidification processes are widely used in the management of hazardous
91 waste.

92 When the waste has been treated, the efficiency of the treatment process must be evaluated, analyzing
93 the untreated and treated waste by chemical analysis, or the so-called leachate tests, which consist of
94 putting the sample in contact with a "leaching" agent during a certain period of time. In this way, the
95 total content of the waste can be compared with that of the treated matrix and what is actually released
96 in contact with the leachant. Let's see what materials make up the study.

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98

99 **2.Materials**

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101 In this section, the materials used for the development of the experiment are analyzed, these
102 being the following.

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104 **2.1 Ferritic Fume Dust (FFD).**

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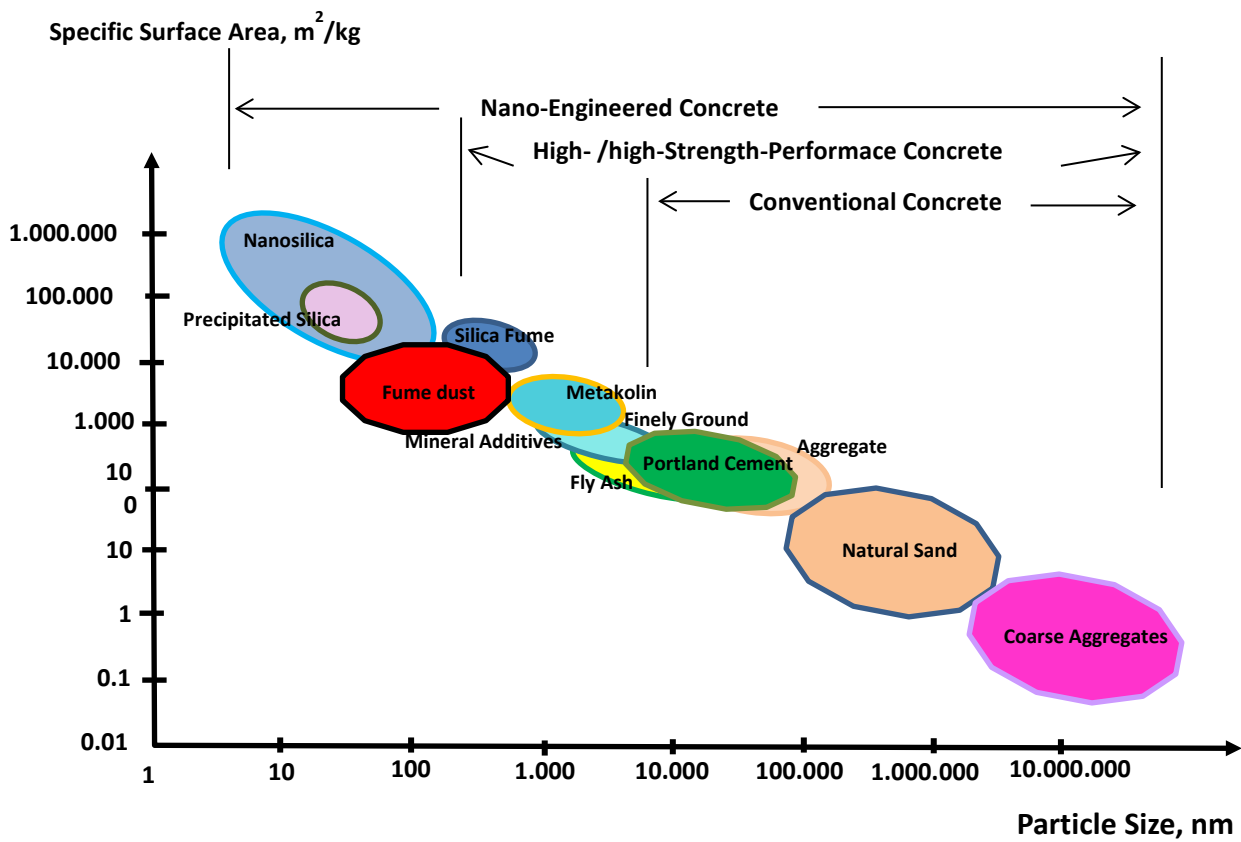
106 Most of our steel industry uses collection and purification systems. The collection system is
107 mainly used in the aspiration through a fourth hole in the roof of the furnace, and subsequent
108 uptake by extractor bells installed in various parts of the mill, mainly above the ovens. The gases
109 captured by the hood are conducted through channels to the fourth filter or bag filters. Emissions
110 generated in the AOD converter (Decarburization converter with Oxygen and Argon) are also
111 captured and directed to the channel and filtered together with the gases of the electric furnace.

112 Thus the solid obtained is a very fine dust material with between 15 and 25 kg of dust produced
113 per ton of steel.

114 Morphologically,[18–20] electric arc furnace (EAF) fume dust consists of spheroidal particles
115 of highly variable diameters (Figures 1 and 2), ranging from 50 to 500 microns. Its moisture
116 content is below 1% by weight. The wide distribution of particle sizes is due to both the
117 composition and origin of the raw materials, as well as differences in the production process.
118 As shown in Figure 1, EAF fume dust, despite being an industrial waste, it has the potential to
119 modify the gel structure, and fill the gaps between the grains of cement and various aggregates
120 increasing hydration of the cement and reducing the capillary void volume of the concrete[21].
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142 *Figure 1. Scale of different types of components*

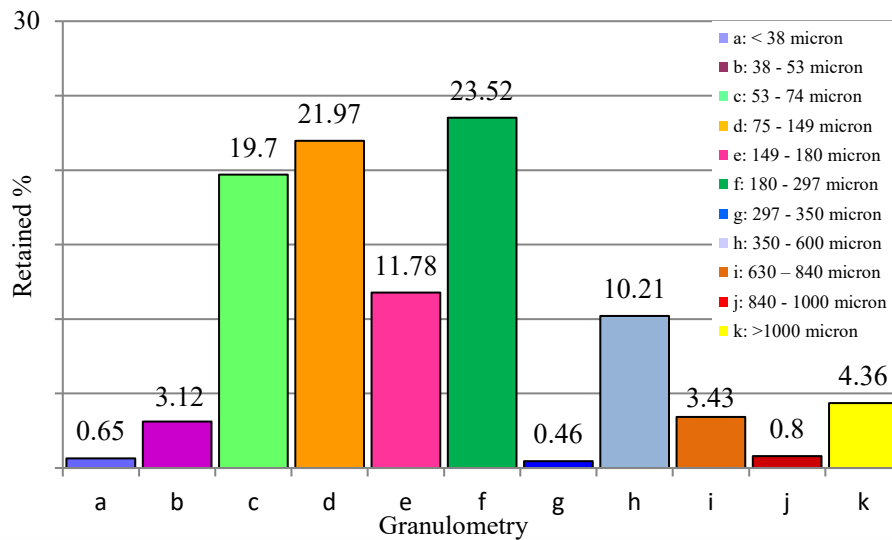
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Figure 2. Ferritic fume dust

157 The sample used in this study was obtained from a random fraction contained in the bag filters where
 158 the waste is stored. From each of these samples approximately 50 kg were taken, always preserved in
 159 closed containers to prevent the action of moisture. Then, 250 g of both samples were taken to proceed
 160 to perform a granulometric study[22].

161 Figure 3 shows the results obtained where the amount of material that does not cross the corresponding
 162 sieve are collected. It is noted that the majority of material is in the range 53-297 microns.



177 *Figure 3 Ferritic fume dust granulometry*

179 The 77.06% of the grains are within this range. The particle size distribution is not symmetrically
 180 shaped but right correspond one 19.17% and to the left a 3.77% of the material analysed[23–25].

181 For the determination of the chemical and mineralogical composition of the EAF fume dust, X-ray
 182 fluorescence, X-ray diffraction (XRD) and thermogravimetric analysis (TGA) were carried out.

183
 184 Table 1 shows the chemical composition obtained by XRF.

Chemical elements	Content %
Carbon (C)	0.20-0.50
Silicon (Si)	2.00-4.00
Manganese (Mn)	1.50-2.50
Tin (Ti)	<0.010
Nickel(Ni)	<0.30
Copper (Cu)	0.20-0.50
Chrome(Cr)	7.00-9.00
Molybdenum(Mb)	0.10-0.20

Iron(Fe)	23.00-32.00
Calcium(Ca)	6.50-8.00
Lead (Pb)	1.00-1.50
Zinc (Zn)	10.00-20.00
Magnesium (Mg)	2.00-3.00

Table 1. Chemical composition of ferritic fume dust obtained by (XRF)

The figure 4 shows the results of thermogravimetric analysis for EAF fume dust sample.

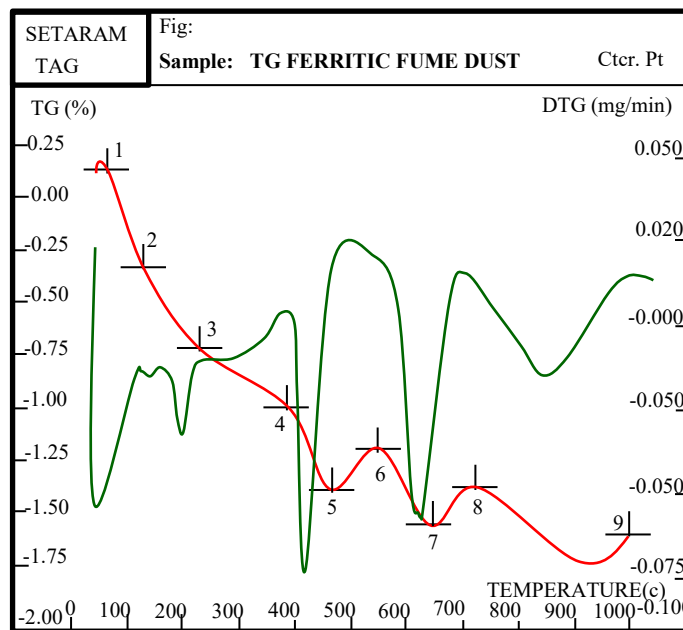


Figure 4 Thermogravimetric analysis of EAF fume dust

Figure 4 shows the thermogravimetric analysis (in red) and differential thermal analysis (shown in green). Figure 5 shows the X-ray diffraction analysis of the sample.

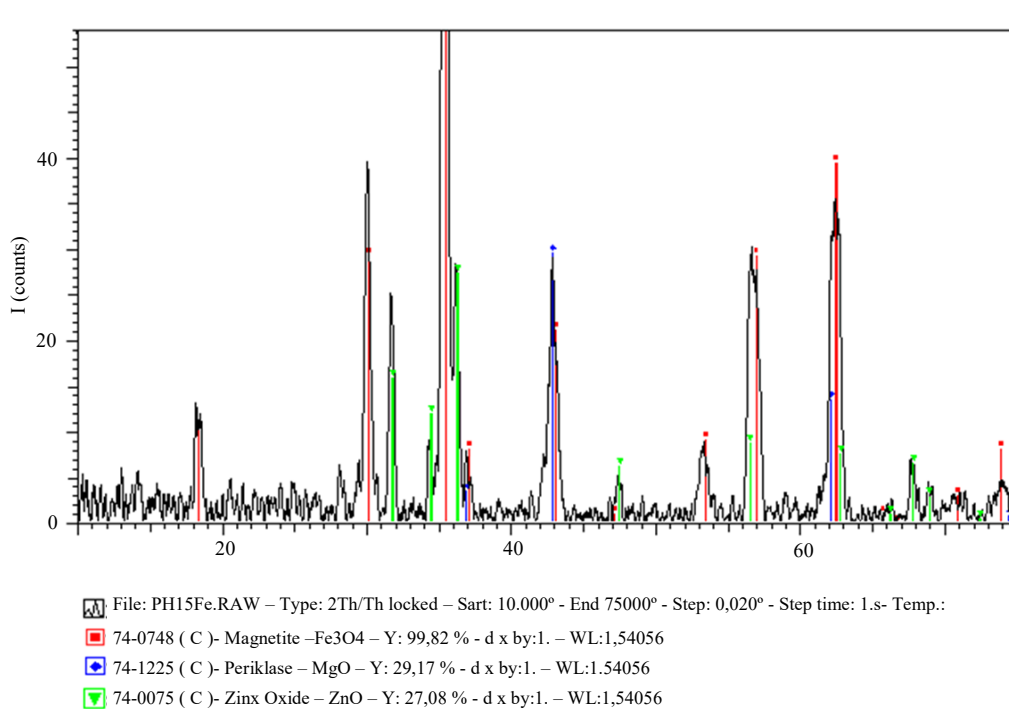


Figure 5 X-ray powder diffraction of EAF fume dust

Powders from the manufacture of stainless steel, consist mainly of magnetite and to a lesser extent, hematite as seen in Figure 5. Calcium and silicon oxides are also present in trace amounts.

Thermal analysis shows an initial zone of continuous loss of mass assignable in principle to dewatering processes commonly occurring in materials having such a fine particle size. However, this loss of absorbed / water constituent peak coincides with the first carbon graphics and humidity obtained by the determiner. This leads us to believe that in this area (below 400 °C) a decomposition product from carbon compounds is also produced along with the water, possibly attributable to organic carbon from oil products. Around 400°C a marked loss occurs due probably to dehydration processes of free calcium hydroxide, which coincides with the second peak of the graph determiner water.

However, in the thermogram, this process is interrupted at 450 °C by a gain region whose origin could be presumed to be oxidation phenomena associated with some kind of structural transformations. One possible structural change to consider is conversion from magnetite to maghemite, according to the reaction



This phenomenon repeats itself later around 600 °C, and likely corresponds to a second transformation of maghemite to hematite according to



Both phenomena of structural transformation (transition from one phase to another) mask losses due to decomposition of carbon and calcium (about 600 °C) in the thermogram, seen in the second peak of

257 the graph, which gives information from the decomposition of alkali carbonates is at temperatures
258 around 800 °C.

259 The actual particle density (3.76 g/cm³) and bulk density (1.06 g/cm³) of the powder sample were
260 obtained using a pycnometer to subsequently make a comparison of densities in the compact.

261

262 2.2. Ordinary Portland Cement

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264 The cement used as binder for every concrete mixture is an ordinary clinker Portland cement supplied
265 by a local company and categorized as type CEM I 52.5 R with a bulk density of 3.1 g/cm³, whose
266 main components are, calcium oxide, silicon oxide, aluminum oxide, iron oxide and magnesium oxide
267 in different proportions according to the manufacturer.

268

269 2.3. Silica Fume

270 Silica fume is an established material for use in high and ultra-high performance concrete. It
271 contributes to strength through filler effects as well as by pozzolanic reaction. Since it falls within a
272 similar size range to the EAF fume dust, it was used for comparative purposes. The properties of the
273 silica fume used in the study are given in Table 2.

274

Appearance	Gray powder
Density at 20 ° C (real)	>2.3 g/cm ³
Apparent density	Aprox. 0,2 g/cm ³
content SiO ₂	> 90%
Chloride content	< 0.1 %

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Table 2. Properties of Silica Fume

277 2.4. Aggregates.

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279 The aggregates used were crushed limestone aggregates, from commercial manufacturing plants
280 located in the Campo de Gibraltar. The proportion of aggregates used in this work was 50% 0-2, 0-4
281 mm sand and 50% 0-16 mm gravel. As shown in figure 6, of the granulometry of each of the aggregates
282 used, according to UNE-EN 933-2.

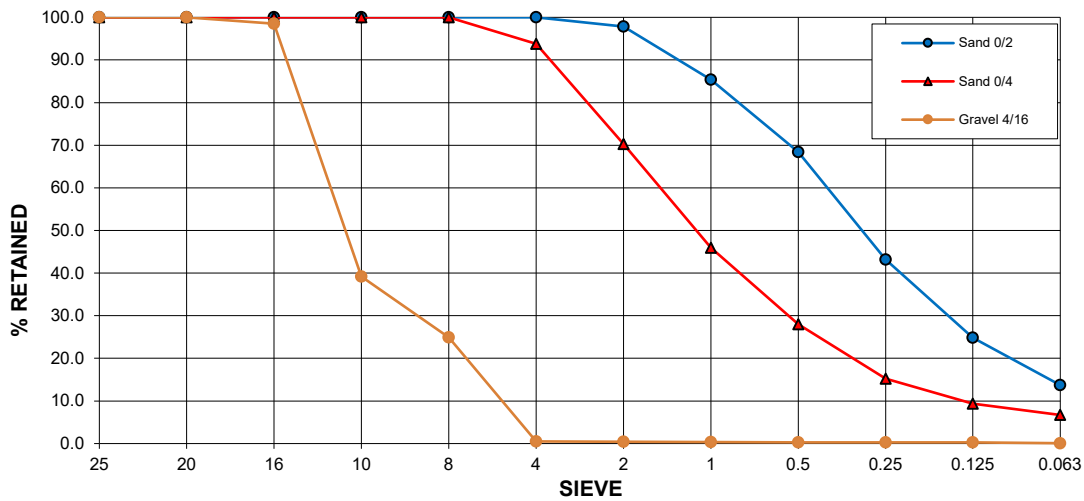


Figure 6 Granulometric sand and gravel

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2.5. Additive.

In order to meet workability requirements of the concrete a superplasticizer additive was used. The additive used in all the concretes has been the Glenium ACE-324, of the commercial mark BASF.

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2.6. Water.

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Domestic tap water was used

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3. Process description

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3.1. Mixture design and manufacture.

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The process for the manufacture of concrete is described below according to the different proportions or values depending on the additions. Six concrete mixtures were produced. Ferritic Fume Dust was included as an addition at levels of 5, 10 and 15%. Two concrete mixtures containing silica fume (10 and 15%) as well as a control mix containing only CEM I were also produced.

306

The nomenclature used for the concrete produced in the study is as follows:

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- FFD 5 = concrete with an addition of 5% (spc) ferritic fume dust.
- FFD 10 = concrete with an addition of 10% (spc) ferritic fume dust.
- FFD 15 = concrete with an addition of 15% (spc) ferritic fume dust.
- SF 10 = concrete with an addition of 10% (spc) silica fume.
- SF 15 = concrete with an addition of 15% (spc) silica fume.

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314 • NA = conventional concrete without addition.

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316 The W/C that has been chosen, in all cases, was 0.5 as we can see in Table 3. The addition of both
317 ferritic smoke dust and silica smoke in small proportions according to UNE-83460-2.

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Mix	Water (w/c ratio)	Binder			Aggregates		
		Cement Dosage Kg/m ³	Addition %	Additive %	Dosage Kg/m ³	Sand 0-4 %	Gravel 4-16 %
NA			0%				
FFD 5			5% of FFD				
FFD 10			10% of FFD				
FFD 15	0.5	325 Kg/m ³	15% of FFD	1.2%	2033.8 kg/m ³	50%	50%
SF 10			10% of SF				
SF 15			15% of SF				

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Table 3. Concrete mixture proportion.

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322 These concretes were manufactured in a mixer with rotating vertical axis and fixed blades, with a
323 capacity of 80 litres. A soft consistency of 7.1 was obtained according to the UNE-EN 12350-2
324 standard.

325 Once mixed the concretes were poured into die and vibrated for compaction, using a vibrating table at
326 a frequency of 42 Hz (2400 cycles per minute) according to UNE 12390-2[26].

327 Specimens were covered with plastic for 24 hours and then were demoulded and brought into a moist
328 chamber for curing at a relative humidity not less than 95% and a temperature of 20 ± 2 ° C. Finally
329 the samples were extruded according to norm UNE-EN 12390-4.

330 These specimens were kept in the chamber until use, 24 hours before testing.

331

332 3.2. Compression strength tests.

333 Compression testing was performed according to standard UNE-EN 12390-3 [27]:

334 The geometry of the test pieces for compression tests were cylindrical according to standard
335 dimensions of diameter d and height $2d$. In our case their dimensions correspond with 45 mm diameter
336 and 90 mm height as shown in Figure 7. According to UNE-EN 12504-1 standard if the thick aggregate
337 does not exceed 20 mm in diameter. Cylindrical specimens were cut without alteration of mortar and
338 coarse aggregate. As the surface was flat, the facing of the test surfaces was dispensed with.

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347 *Figure 7 (a) Shows a cylindrical specimen.*

Figure 7 (b) Sample prismatic specimen.

348 For performing compression tests, three replicate specimens for each concrete mix were tested at ages
349 of 7, 28 and 90 days.

351 3.3 Flexural strength tests.

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353 The flexural test samples with the same characteristics were performed using prismatic specimens with
354 dimensions according to regulations square section edge d and length $2d$ or $4d$, in our case the
355 dimensions of these specimens are 40 x 40 x 160 mm as shown in Figure 8, according to UNE 12390-
356 1 standard, provided that $L \geq 3.5 d$ in this case $3.5 \times 16 = 56$ mm.

357 Following the flexural test, the two halves originating from the failed specimen were tested in
358 compression. As the section is square, one can obtain a modified or equivalent cube load applied by
359 square plates with the same dimensions as the cross section of the prism.



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369 *Figure 8. Charging device cubic specimens for compressive*

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371 According to Neville [5,14,17,28] the resistance of the modified specimen, would be 5% higher than
372 the normal cube specimen of the same size, because of the lateral containment due to excesses in
373 relation to the hub. In this study an average relationship is obtained, since the test was performed using
374 2 for each of the specimens tested for flexural strength. Said test is carried out as a comparative of the
375 previous one.

377 3.4 Leaching test.

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379 The study was developed based on the UNE-EN 12920 standard. The different processes and steps to
380 be followed have been applied according to this regulation.

381 This standard provides a methodology to determine the leaching behavior of a waste under specific
382 conditions, that is, in a solidification / stabilization scenario within a specific time frame, in our case
383 corresponding to 100 years. A selection of tests is required depending on the problem and the scenario
384 to be evaluated [29,30].

385 This methodology is specific to determine a leachate behavior of a waste under specific conditions,
386 since the stabilization / solidification of the material is used. Therefore, external conditions that have
387 an influence on the release of components of the waste in question were considered.

388 It is intended to see if said matrix is capable of retaining the heavy metals within it without any
389 deterioration in the material being produced, under accelerated conditions and subsequently
390 the determination of the leaching behavior within the specified time frame.

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392 To determine the behavior in the material, two different tests were carried out,

393 • The first was to study specimens of 45mm in diameter and 90 mm in height, with an addition
394 of 15% FFD, this being the highest amount of fume dust considered in this research. These were
395 cured for 28 days in wet camera to see the proportions of heavy materials that gave off in this
396 first leaching.

397

398 The test sample was immersed in 800 ml of distilled water for 24 hours. After the test period, the
399 sample was extracted and the leachate solution was transferred to a 1000 mL volumetric flask and
400 made up to the mark with distilled water. The test was repeated by immersing the test sample for 24
401 hours more in another 800mL of water, repeating the described methodology.

402

403 • In the second test, the specimens were aged rapidly. After 28 days of curing in a humid
404 chamber, test samples were subjected to oven drying at 60°C for 48 hours and then immersed in
405 a water bath at 70 C for another 32 days. Then they were deposited on the street exposed to the
406 elements for 3 months. Once aged, the same technique described in the previous paragraph for
407 the leaching test was developed.

408

409 **4. Results and discussion**

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411 4.1 Flexural and compressive strength

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413 In Table 4 the results of tests with cylindrical specimens are detailed. The table shows the average
414 values obtained after breaking three specimens in each case, being the evaluation of rupture mode,
415 according to regulations, satisfactory.

416

Mix description	7 days (MPa)	28 days (MPa)	90 days (MPa)
FFD 5	38.3	41.7	45.3
FFD 10	41.3	40.2	54.8
FFD 15	50.6	51.6	61.8
SF 10	43.1	46.2	59.5
SF 15	38.3	42.2	52.5
NA	37.3	39.1	51.3

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Table 4. Results of compression tests.

Following the flexural test, the two halves originating from the tested specimen were tested in compression. Results of flexural strength test at 28 days and compressive strength at 40 days are shown in Table 5 and Table 6 respectively.

Sample	28 days (MPa)
FFD 5	13.9
FFD 10	12.6
FFD 15	11.1
SF 10	11.9
SF 15	11.1
NA	8.65

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Table 5 Results of prismatic specimens, flexural tests.

Mixed description(%)	Break to 40 days (MPa)
FFD 5	55.71
FFD 10	54.80
FFD 15	70.25
SF 10	69.87
SF 15	57.54
NA	57.44

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Table 6. Results of cubic specimens. Compression tests.

In various tests, particularly for cylindrical samples, compressive strength increased with increasing amount of powder of ferritic fume dust in the concrete composition, while the addition of silica fume decreases effect and Kadri referenced by Duval[31,32] .

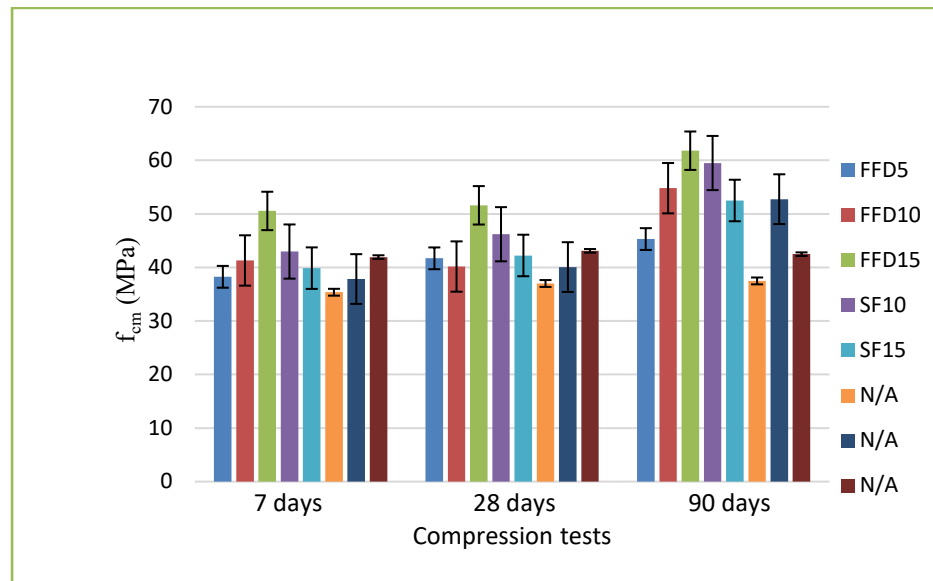


Figure 9. Compressive strength on cylindrical specimens (error bars indicate the standard deviation).

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440 To 7 days appreciate that increasing the strength of the material begins to be significant primarily with
441 the addition of 15% fume dust as surpasses even high strength concrete made adding silica fume.
442 At 28 days the increase is quite relevant, compared to the rest, when the addition is 15% ferritic fume
443 dust.

444 90 days note that unlike what happens with silica fume powder, the higher the addition of powdered
445 ferritic fume dust, compression strength of the material increases, reaching a difference of 9.3 MPa
446 with respect SF15% of more than 15% increase, although with FFD10% is 2,3MPa only slightly less
447 than 5% increase. Draw attention to the strong improvement produced with the addition of fume dust
448 respect to silica fume.

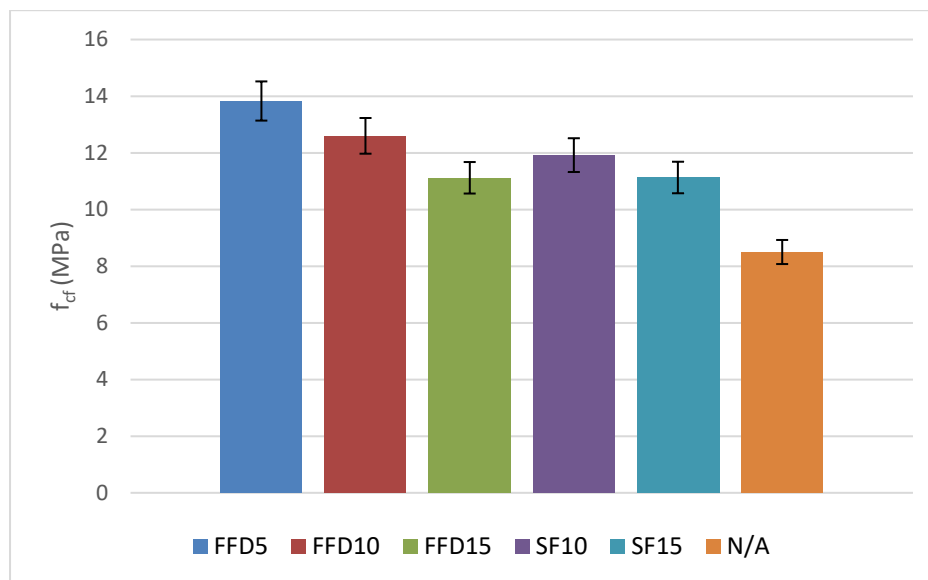
449 If the FFD15% compared with conventional concrete gain it is 10.5 MPa, up from 15% the difference
450 with NA.

451 The results show that concrete with 15% ferritic fume dust (FFD15) acquire a higher resistance again
452 in respect of comparative equivalent performed for cylindrical specimens for both concretes with
453 added silica fume 15 % and 10% and conventional concrete. That is, FFD15 increased by 2% over the
454 SF10, up 18% compared to SF10 equal increment on concrete NA.

455 Contrary to the results obtained for compression, and as shown in Table 6, in the case of the flexural
456 strength, are higher values as decreases powder addition ferritic fume dust, jumping almost 14 MPa of
457 FFD5 the value of 11 MPa for the addition FFD15. This is a loss of about 25% of the addition as we
458 increase resistance of ferritic fume dust. This fact must be taken into account as previously exhibited
459 levels of fissuring obtained by the addition, resulting in a positive development since the flexural
460 strength obtained by adding PHF15 is similar to that obtained for concrete with addition powder silica
461 fume, then the levels of cracking in the absence of a formal verification is not the subject of this work,
462 will be equivalent.

463 If we make a comparison with conventional concrete, increased flexural strength is notable for each
464 addition FFD15, FFD10 and FFD5 rising from just over 25% higher than Portland cement concrete to

465 almost 60%, respectively. This data supports the idea of the decrease in cracking of concrete with
 466 added ferritic fume dust.
 467 In Figure 10 the bar chart with this line down on resistance just discussed is located.
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469
 470 *Figure 10. Flexural strength of concrete (error bars indicate the standard deviation).*
 471

472 FFD and SF has a similar behaviour as nano particles that fill the voids of the cement matrix [21,33].
 473 For this reason, an improvement of the mechanical characteristics of the concrete mixtures studied in
 474 this work has been observed.

475
 476 **4.2 Leaching analysis**
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478 Tables 7, 8 and 9 show the results corresponding to the total content of heavy and non-heavy metals.
 479 The results are shown in parts per million (ppm). In the different tables a comparison is made between
 480 the simply cured and aged samples. The weights of the test pieces are taken in grams for the
 481 approximation in mg / kg of dry matter.
 482

Components Test tubes			Ferritic fume dust (FFD 5%)		Conventional concrete	
Mass Test tubes g			365.95		353,00	
Leaching			1st Non-aged leaching	2nd Non-aged leaching	1st Non-aged leaching	2nd Non-aged leaching
Detection limits	0.1	Ca	1.59	0.9	1.56	1.76
	0.004	Cr _{Total}	0.045	0.057	0.007	0.023
	0.042	Fe	0.062	<LD	<LD	<LD
	0.09	SO ₄ ⁼	6.91	2.17	17.90	14.15
	0.014	Zn	0.016	0.002	0.014	<LD

1.9	Na	22.51	10.38	16.81	7.15
0.001	Mn	0.001	<LD	<LD	<LD
0.014	Al	0.095	0.066	0.071	0.042
0.056	K	37.27	18.62	28.15	17.87
0.025	Si	2.18	1.76	2.75	1.190
TDS (ppm)		101	57.46	82.59	54.09
Conductivity (μ S / cm)		179	103.5	147.6	97.6
pH		9.250	8.680	8.990	8.160

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Table 7 Ferritic fume dust leaching result (FFD5%) results with conventional concrete in non-aged specimens

Components Test tubes		Ferritic fume dust (FFD10)		Conventional concrete		
Mass Test tubes g		365.95		353.26		
Leaching		1st Non-aged leaching	2nd Non-aged leaching	1st Non-aged leaching	2nd Non-aged leaching	
Detection limits (PPM)	0.1	Ca	1.59	0.95	1.56	1.76
	0.004	Cr _{Total}	0.046	0.067	0.007	0.024
	0.042	Fe	0.071	<LD	<LD	<LD
	0.001	Mn	0.001	<LD	<LD	<LD
	0.09	SO ₄	10.09	2.78	17.90	14.15
	0.002	Zn	0.014	<LD	0.014	<LD
	1.9	Na	24.15	10.22	16.81	7.15
	0.014	Al	0.084	0.066	0.071	0.042
	0.056	K	48.46	22.88	28.15	17.87
	0.025	Si	2.48	1.89	2.75	1.90
TDS (ppm)		117	63.67	82.59	54.09	
Conductivity (μ S / cm)		215	114.4	147.6	97.6	
pH		9.3	8.6	8.990	8.160	

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Table 8 Ferritic fume dust leaching result (FFD10%) results with conventional concrete in non-aged specimens .

Components Test tubes		Ferritic fume dust (FFD15)		Conventional concrete	
Mass Test tubes g		362.00		353.26	
Leaching		1st Non-aged leaching	2nd Non-aged leaching	1st Non-aged leaching	2nd Non-aged leaching

Detection limits (PPM)	0.1	Ca	1.971	2.603	1.56	1.76
	0.004	Cr _{Total}	0.005	<LD	0.007	0.023
	0.09	SO ₄	<LD	<LD	17.90	14.15
	0.002	Zn	<LD	<LD	0.014	<LD
	1.9	Na	<LD	<LD	16.81	7.15
	0.014	Al	<LD	<LD	0.071	0.042
	0.056	K	1.806	0.978	28.15	17.87
	0.025	Si	0.228	0.240	2.75	1.90
TDS (ppm)			13.87	11.91	82.59	54.09
Conductivity (μS / cm)			27.04	23.6	147.6	97.6
pH			8.61	8.15	8.99	8.16

Table 9 Ferritic fume dust leaching result (FFD15%) results with conventional concrete in non-aged specimens

From the observation in Table 7 an immediate conclusion is drawn, the parameters that are found in greater abundance in the first leaching is the value of the Cr_{Total} although lower in the first balance due to the decrease in PH and the consequent partial dilution, since in the 2nd leaching it grows instead of decreasing. As expected in the second leaching, there is generally a decrease in the concentration of leached species[34,35].

Also note that in Tables 8 and 9 the amounts of total chromium in conventional concrete samples are greater than those given for concrete added with ferritic fume dust.

On page 36 of Directive [36,37] Official Journal of the European Communities DO L 11 de 16.1.2003, p. 27/48; Directive with the Council Decision of 19/12/2002 2003/33/CE on "Criterion and acceptance procedures waste in landfills". The maximum limit for total chromium is 4 mg / Kg of dry matter calculated in terms of total release, for proportions between liquid and solid (L/S) of 2 L/kg. This value is the least permissive of the set of tables that are dictated in this European Directive. If we make the balance in mg / kg of dry matter of the total chromium of table 9 above and with an easy proportion, it is determined that in our samples the leaching is lower

$$\frac{0,057 \text{ mg} \times 1000 \text{ g}}{362,41 \text{ g}} \Rightarrow 0,157 \frac{\text{mg}}{\text{kg}} \text{ dry materials}$$

[3]

Comparing it with a liter of water, if we take it to the proportion formulated in the table of 2 L/kg, this value reaches 0.315 mg/kg well below the maximum limit of 4 mg/kg imposed by the standard.

514 The variability of other metal oxides is large qualitatively but quantitatively negligible, being narrower
515 marked previously in the Cr representing one tenth of the assigned limit. The average of these
516 parameters is less than 5%.

517 In addition to the leaching limit values mentioned in Table 8, granular residues should meet the
518 following additional criteria:

519

520 pH parameter ----- Limit value ≥ 6

521

522 The lowest value registered in the analytical is 8.15, higher than that indicated in the standard.

523

524 A simple observation of the comparative tables 8 and 9 give value to the performance of another
525 species in leaching. The sulphate parameter decreases in the concrete samples with ferritic fume dust
526 with respect to the amount of this same measure in conventional concrete.

527

528

529 **5. Conclusions**

530 The main findings throughout the work are listed below:

531

- 532 • Ferritic fume dust used as additive to concrete, gives a good workability, compactness and
533 rigidity hardened material.
- 534 • Concrete mixtures with ferritic fume dust shown an improvement of their compressive
535 strength in comparison with concrete mixtures with silica fume (both with the same
536 proportion).
- 537 • The mechanical-resistant behavior of the concrete with the addition of ferritic fume dust
538 powder achieves compressive and flexo-elastic resistances superior to conventional concrete.
- 539 • This material is classified as a special waste because it exceeds the limit of chromium, but this
540 and other contaminants are stabilized in the cement matrix, as we have seen in the leaching of
541 the concrete.
- 542 • It can be considered a micro-filler that can be used as a mineral addition in high or ultra-high
543 performance concrete.
- 544 • A technique of stabilization/solidification of the ferritic fume dust with concrete was chosen,
545 evaluating different proportions of this residue 5%, 10% and 15% of addition.
- 546 • First group not aged: Only the concentration values in leachate in chromium are somewhat
547 higher than those of conventional concrete and very little above zinc and aluminum.
- 548 • Second group aged: Since in the total chrome it becomes less than in conventional concrete.
- 549 • The fact that in some elements such as total chromium that present slightly higher values in the
550 second leaching, is motivated by the decrease in pH and the consequent partial dissolution of
551 its hydroxides.

552

553

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555 analyzed the results; M.D.R.-C., S.B, F.P.-G. and M.E.P.-R. performed the experiments.

556

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559

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561 **Declaration statement:** The data that support the findings of this study are available from the
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